WEST Search History

Hide Items | Restore | Clear | Cancel

DATE: Tuesday, December 05, 2006

Hide?	Set Name	Query	Hit Count
		DB=USPT; PLUR=YES; OP=ADJ	
☐	L2	L1 and piperidin$3	276
☐	L1	(514/$7.ccls or 544/$7.ccls. or 546/$7.ccls.) and phospholipase	400

END OF SEARCH HISTORY

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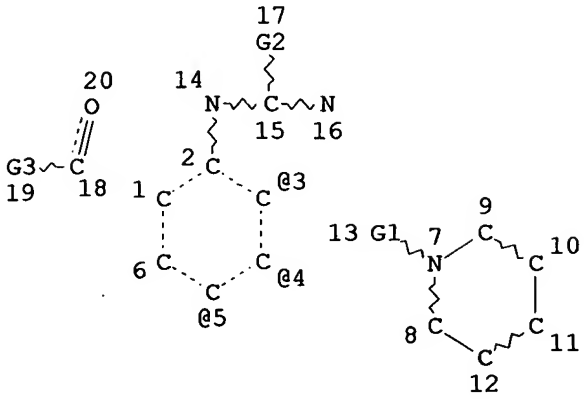
L4 HAS NO ANSWERS
L4 STR



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VAR G2=O/S
VAR G3=N/CY
NODE ATTRIBUTES:
DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:
RSPEC 8 1
NUMBER OF NODES IS 20

STEREO ATTRIBUTES: NONE

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(FILE 'REGISTRY' ENTERED AT 11:56:54 ON 05 DEC 2006)
L6 1888 S L4 FUL

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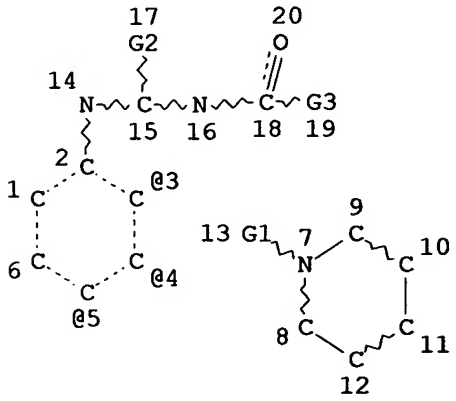
L8 HAS NO ANSWERS
L8 STR



VAR G1=3/4/5
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VAR G3=N/CY
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DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:
RSPEC 8 2
NUMBER OF NODES IS 20

STEREO ATTRIBUTES: NONE

=> search l8
ENTER TYPE OF SEARCH (SSS), CSS, FAMILY, OR EXACT:sss
ENTER SCOPE OF SEARCH (SAMPLE), FULL, RANGE, OR SUBSET:subset
ENTER SUBSET L# OR (END):16
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FULL SUBSET SEARCH INITIATED 12:06:53 FILE 'REGISTRY'
FULL SUBSET SCREEN SEARCH COMPLETED - 756 TO ITERATE

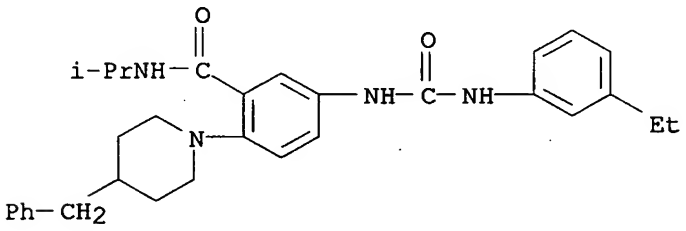
100.0% PROCESSED 756 ITERATIONS 756 ANSWERS
SEARCH TIME: 00.00.01

L9 756 SEA SUB=L6 SSS FUL L8

=> s l6 not l9
L10 1132 L6 NOT L9

=> d scan

L10 1132 ANSWERS REGISTRY COPYRIGHT 2006 ACS on STN
IN Benzamide, 5-[[[(3-ethylphenyl)amino]carbonyl]amino]-N-(1-methylethyl)-2-[4-(phenylmethyl)-1-piperidinyl]- (9CI)
MF C31 H38 N4 O2



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):0

=> fil caplus
COST IN U.S. DOLLARS SINCE FILE ENTRY TOTAL SESSION
FULL ESTIMATED COST 41.16 438.76

FILE 'CAPLUS' ENTERED AT 12:07:14 ON 05 DEC 2006
USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.
PLEASE SEE "HELP USAGETERMS" FOR DETAILS.

FILE COVERS 1907 - 5 Dec 2006 VOL 145 ISS 24
FILE LAST UPDATED: 4 Dec 2006 (20061204/ED)

Effective October 17, 2005, revised CAS Information Use Policies apply. They are available for your review at:

http://www.cas.org/infopolicy.html

=> s l10
L11 37 L10

=> d bib 1-37

L11 ANSWER 1 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2006:485850 CAPLUS
DN 144:495337
TI Substituted biaryl-carboxylate derivatives as bradykinin B1 antagonists or inverse agonists useful in the treatment of pain and inflammation
IN Wood, Michael R.; Bock, Mark G.; Books, Kathy M.; Freidinger, Roger M.; Kim, June J.
PA USA
SO U.S. Pat. Appl. Publ., 18 pp.
CODEN: USXXCO
DT Patent
LA English
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI US 2006111392	A1	20060525	US 2005-284740	20051122
PRAI US 2004-630594P	P	20041123		

OS MARPAT 144:495337

L11 ANSWER 2 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2006:164439 CAPLUS
DN 144:253908
TI Preparation of aryl urea derivatives as CB1 cannabinoid receptor modulators
IN Bloxham, Jason; Fyfe, Matthew Colin Thor; Horswill, James; Jeevaratnam, Revathy Perpetua; Keily, John; Procter, Martin James; Schofield, Karen Lesley; Shaaban, Salam; Swain, Simon Andrew; Wong-Kai-In, Philippe
PA Prosidion Limited, UK
SO PCT Int. Appl., 66 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2006018662	A2	20060223	WO 2005-GB50131	20050816

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA,

NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK,
SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU,
ZA, ZM, ZW
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE,
IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ,
CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH,
GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
KG, KZ, MD, RU, TJ, TM
PRAI US 2004-602268P P 20040816
OS MARPAT 144:253908

L11 ANSWER 3 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2006:164356 CAPLUS
DN 144:254385
TI Preparation of tripeptides bearing a cyclopropyl moiety and phosphorous groups as antiviral compounds
IN Chaudhary, Kleem; Fleury, Melissa; Kim, Choung U.; Mcmurtrie, Darren J.; Sheng, Xiaoning C.
PA Gilead Sciences, Inc., USA
SO PCT Int. Appl., 476 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2006020276	A2	20060223	WO 2005-US25503	20050718
WO 2006020276	A3	20061005		

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH,
CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD,
GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ,
LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA,
NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK,
SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU,
ZA, ZM, ZW
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE,
IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ,
CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH,
GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
KG, KZ, MD, RU, TJ, TM

US 2006122123	A1	20060608	US 2005-184429	20050718
PRAI US 2004-588633P	P	20040716		
US 2004-591635P	P	20040727		

OS MARPAT 144:254385

L11 ANSWER 4 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2006:31771 CAPLUS
DN 144:129003
TI Preparation of urea derivatives as acyl-CoA:diacylglycerol acyltransferase (DGAT) inhibitors
IN Kurata, Hitoshi; Uto, Yoshikazu; Fujibayashi, Yuko; Kohama, Takafumi; Tanimoto, Tatsuo; Karasawa, Hiroshi
PA Sankyo Company, Limited, Japan
SO PCT Int. Appl., 524 pp.
CODEN: PIXXD2
DT Patent
LA Japanese
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2006004200	A1	20060112	WO 2005-JP12635	20050701

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH,

CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW

RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

JP 2006045209 A2 20060216 JP 2005-193260 20050701

PRAI JP 2004-196723 A 20040702

OS MARPAT 144:129003

RE.CNT 10 THERE ARE 10 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 5 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN

AN 2005:1039511 CAPLUS

DN 145:145355

TI Liquid-phase parallel synthesis of substituted 4-aminobenzamides

AU Trifilenkov, A. S.; Il'in, A. P.; Kravchenko, D. V.; Dorogov, M. V.; Blyumina, M. V.; Ivashchenko, A. V.

CS Yarosl. Gos. Pedagog. Univ. im. K. D. Ushinskogo, Yaroslavl, Russia

SO Izvestiya Vysshikh Uchebnykh Zavedenii, Khimiya i Khimicheskaya Tekhnologiya (2005), 48(5), 137-144
CODEN: IVUKAR; ISSN: 0579-2991

PB Ivanovskii Gosudarstvennyi Khimiko-Tekhnologicheskii Universitet

DT Journal

LA Russian

OS CASREACT 145:145355

L11 ANSWER 6 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN

AN 2005:977019 CAPLUS

DN 143:286162

TI Preparation of aryl semicarbazide derivatives as kinase inhibitors

IN Buchstaller, Hans-Peter; Finsinger, Dirk; Stieber, Frank; Wiesner, Matthias; Amendt, Christiane; Sirrenberg, Christian; Zenke, Frank; Grell, Matthias

PA Merck Patent G.m.b.H., Germany

SO PCT Int. Appl., 278 pp.
CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2005082853	A1	20050909	WO 2005-EP1443	20050214

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW

RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

AU 2005217041	A1	20050909	AU 2005-217041	20050214
CA 2557359	AA	20050909	CA 2005-2557359	20050214

PRAI EP 2004-4330 A 20040226

WO 2005-EP1443 W 20050214
OS MARPAT 143:286162
RE.CNT 11 THERE ARE 11 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 7 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2005:732627 CAPLUS
DN 143:211919
TI Preparation of heterocyclic urea derivatives as coagulation factor Xa inhibitors.
IN Mederski, Werner; Tsaklakidis, Christos; Dorsch, Dieter; Cezanne, Bertram; Gleitz, Johannes
PA Merck Patent G.m.b.H., Germany
SO PCT Int. Appl., 57 pp.
CODEN: PIXXD2
DT Patent
LA German
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2005073201	A1	20050811	WO 2005-EP83	20050107
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
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DE 102004004731	A1	20050818	DE 2004-102004004731	20040130
AU 2005209362	A1	20050811	AU 2005-209362	20050107
CA 2554911	AA	20050811	CA 2005-2554911	20050107
EP 1709017	A1	20061011	EP 2005-700739	20050107
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, CY, TR, BG, CZ, EE, HU, PL, SK, IS				

PRAI DE 2004-102004004731 A 20040130
WO 2005-EP83 W 20050107
OS CASREACT 143:211919; MARPAT 143:211919
RE.CNT 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 8 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2004:1127325 CAPLUS
DN 142:74359
TI Synthesis of N-hydroxy-7-(arylamino)heptanamide derivatives useful for treating hyper-proliferative disorders
IN Kluender, Harold C. E.; Hong, Zhenqiu; Ladouceur, Gaetan H.; Liu, Xiao-Gao; Khire, Uday; Wang, Lei
PA Bayer Pharmaceuticals Corporation, USA
SO PCT Int. Appl., 82 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2004110989	A1	20041223	WO 2004-US15465	20040513
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD,				

GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,
LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI,
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TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
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AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK,
EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE,
SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE,
SN, TD, TG
PRAI US 2003-470713P P 20030514
OS MARPAT 142:74359
RE.CNT 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 9 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2004:1080898 CAPLUS
DN 142:56358
TI Preparation of aroylsemicarbazides as factor Xa inhibitors for the treatment of thromboembolic diseases
IN Mederski, Werner; Tsaklakidis, Christos; Dorsch, Dieter; Cezanne, Bertram; Gleitz, Johannes
PA Merck Patent G.m.b.H., Germany
SO PCT Int. Appl., 36 pp.
CODEN: PIXXD2
DT Patent
LA German
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2004108718	A1	20041216	WO 2004-EP5088	20040512
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
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DE 10325962	A1	20041223	DE 2003-10325962	20030607
AU 2004245187	A1	20041216	AU 2004-245187	20040512
CA 2528233	AA	20041216	CA 2004-2528233	20040512
EP 1633745	A1	20060315	EP 2004-732283	20040512
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, CY, TR, BG, CZ, EE, HU, PL, SK				
BR 2004010617	A	20060620	BR 2004-10617	20040512
CN 1802370	A	20060712	CN 2004-80015854	20040512
JP 2006527217	T2	20061130	JP 2006-515768	20040512
US 2006241111	A1	20061026	US 2006-559385	20060621
PRAI DE 2003-10325962	A	20030607		
WO 2004-EP5088	W	20040512		

RE.CNT 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 10 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2004:878375 CAPLUS
DN 141:350047
TI Preparation of phospholipase C inhibitors for use in treating inflammatory diseases
IN Lagu, Bharat; Rupert, Kenneth; Wachter, Michael

PA Janssen Pharmaceutica N.V., Belg.
SO PCT Int. Appl., 114 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2004089901	A2	20041021	WO 2004-US9847	20040331
WO 2004089901	A3	20041209		

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RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

US 2004242639	A1	20041202	US 2004-814070	20040331
PRAI US 2003-459078P	P	20030331		

OS MARPAT 141:350047

L11 ANSWER 11 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2004:878168 CAPLUS
DN 141:360665
TI Synergistic methods and compositions using insulin-like growth factor 1 receptor (IGF1R) inhibitors with additional kinase inhibitors for treating cancer
IN Carboni, Joan M.; Hurlburt, Warren W.; Gottardis, Marco M.; Lee, Francis Y.
PA USA
SO U.S. Pat. Appl. Publ., 66 pp., Cont.-in-part of U.S. Ser. No. 676,214.
CODEN: USXXCO
DT Patent
LA English
FAN.CNT 3

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI US 2004209930	A1	20041021	US 2004-814199	20040331
CA 2500714	AA	20040415	CA 2003-2500714	20031001
US 2004072760	A1	20040415	US 2003-677067	20031001
AU 2003275364	A1	20040423	AU 2003-275364	20031001
US 2004106605	A1	20040603	US 2003-676214	20031001
EP 1551411	A2	20050713	EP 2003-759640	20031001

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK

JP 2006503867	T2	20060202	JP 2004-541997	20031001
WO 2005094376	A2	20051013	WO 2005-US10820	20050330

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

PRAI US 2002-415416P P 20021002
US 2003-676214 A2 20031001
US 2003-677067 A2 20031001
WO 2003-US31091 W 20031001
US 2004-814199 A 20040331
OS MARPAT 141:360665

L11 ANSWER 12 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2004:857548 CAPLUS
DN 141:350049
TI Preparation of (hetero)arylurea derivatives as deformylase inhibitors with antibacterial activity
IN Lee, Bong-Jin; Lee, Seung-Kyu; Choi, Kwang-Hyun; Lee, Sang-Jae
PA Promeditech Inc., S. Korea
SO PCT Int. Appl., 64 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2004087643	A1	20041014	WO 2004-KR502	20040311

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

KR 2004086171	A	20041008	KR 2004-14549	20040304
PRAI KR 2003-20486	A	20030401		

OS MARPAT 141:350049
RE.CNT 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 13 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2004:696342 CAPLUS
DN 141:225302
TI Preparation of N-arylheterocycles as melanin concentrating hormone (MCH) antagonists.
IN Schwink, Lothar; Stengelin, Siegfried; Gossel, Matthias; Boehme, Thomas; Hessler, Gerhard; Stahl, Petra; Gretzke, Dirk
PA Aventis Pharma Deutschland GmbH, Germany; Aventis Pharma GmbH
SO PCT Int. Appl., 390 pp.
CODEN: PIXXD2
DT Patent
LA German
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2004072025	A2	20040826	WO 2004-EP1342	20040213
WO 2004072025	A3	20041223		

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI
RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU,

MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

	DE 10306250	A1	20040909	DE 2003-10306250	20030214
	AU 2004212145	A1	20040826	AU 2004-212145	20040213
	CA 2516118	AA	20040826	CA 2004-2516118	20040213
	EP 1597228	A2	20051123	EP 2004-710808	20040213
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
	BR 2004007504	A	20060214	BR 2004-7504	20040213
	CN 1774418	A	20060517	CN 2004-80009860	20040213
	JP 2006517563	T2	20060727	JP 2006-501827	20040213
	US 2004220191	A1	20041104	US 2004-779853	20040217
	NO 2005004220	A	20051028	NO 2005-4220	20050912
PRAI	DE 2003-10306250	A	20030214		
	US 2003-488545P	P	20030718		
	WO 2004-EP1342	A	20040213		

OS MARPAT 141:225302

L11 ANSWER 14 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2004:220082 CAPLUS
DN 140:253556
TI Preparation of 5-thiazolecarboxamides as protein tyrosine kinase inhibitors
IN Das, Jagabandhu; Padmanabha, Ramesh; Chen, Ping; Norris, Derek J.; Doweyko, Arthur M. P.; Barrish, Joel C.; Wityak, John; Lombardo, Louis J.; Lee, Francis Y. F.
PA Bristol-Myers Squibb Company, USA
SO U.S. Pat. Appl. Publ., 184 pp., Cont.-in-part of U.S. 6,596,746.
CODEN: USXXCO
DT Patent
LA English
FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 2004054186	A1	20040318	US 2003-395503	20030324
	US 7125875	B2	20061024		
	US 6596746	B1	20030722	US 2000-548929	20000413
	US 2004024208	A1	20040205	US 2003-378372	20030303
	US 6979694	B2	20051227		
	US 2004073026	A1	20040415	US 2003-378461	20030303
	US 7091223	B2	20060815		
	US 2004077875	A1	20040422	US 2003-378373	20030303
	AU 2004223828	A1	20041007	AU 2004-223828	20040323
	CA 2519898	AA	20041007	CA 2004-2519898	20040323
	WO 2004085388	A2	20041007	WO 2004-US8827	20040323
	WO 2004085388	A3	20050630		
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
	RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	EP 1610780	A2	20060104	EP 2004-758053	20040323
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, PL, SK				
	BR 2004008782	A	20060328	BR 2004-8782	20040323
	CN 1764454	A	20060426	CN 2004-80007845	20040323

	JP 2006523216	T2	20061012	JP 2006-507475	20040323
	US 2005261305	A1	20051124	US 2005-138793	20050525
	US 2005288303	A1	20051229	US 2005-138942	20050526
	NO 2005004359	A	20051019	NO 2005-4359	20050920
	US 2006079563	A1	20060413	US 2005-271626	20051110
PRAI	US 1999-129510P	P	19990415		
	US 2000-548929	A2	20000413		
	US 2003-378373	A1	20030303		
	US 2003-395503	A	20030324		
	WO 2004-US8827	W	20040323		

OS MARPAT 140:253556
RE.CNT 149 THERE ARE 149 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 15 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2004:101158 CAPLUS
DN 140:146014
TI Preparation of 4-[[(1-acylaminocyclohexyl)carbonyl]amino]-1-phenylpiperidin-3-ones as cysteine protease inhibitors and processes for their preparation
IN Lee, Jong-Wook; Lee, Bong-Yong; Lee, Chun-Ho; Hur, Yun; Han, Tae-Dong; Ko, Hyun-Kyoung; Yun, Suk-Won; Shim, Jae-Young; Lim, Joong-In; Son, Moon-Ho; Yang, Jae-Sung; Kim, Mi-Kyung
PA Yuhan Corporation, S. Korea; Dong-A Pharmaceutical Co., Ltd.
SO PCT Int. Appl., 158 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2004011457	A1	20040205	WO 2003-KR1502	20030726
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	KR 2004010407	A	20040131	KR 2003-51574	20030725
	AU 2003281698	A1	20040216	AU 2003-281698	20030726
	US 2005234057	A1	20051020	US 2005-521752	20050119
PRAI	KR 2002-44164	A	20020726		
	KR 2003-13889	A	20030306		
	WO 2003-KR1502	W	20030726		

OS MARPAT 140:146014
RE.CNT 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 16 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2003:174468 CAPLUS
DN 138:215278
TI Method of treating hyperresorptive bone disorders by inhibition of Src protein tyrosine kinase
IN Safar, Pavel; Walser, Armin
PA USA
SO U.S. Pat. Appl. Publ., 33 pp.
CODEN: USXXCO
DT Patent

LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 2003045480	A1	20030306	US 2002-191446	20020709
PRAI	US 2001-303851P	P	20010709		

OS MARPAT 138:215278

L11 ANSWER 17 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2003:58070 CAPLUS
DN 138:122861
TI Preparation of substituted amides, sulfonamides and ureas useful for inhibiting kinase activity
IN Safar, Pavel; Walser, Armin; Shimshock, Stephen J.
PA Aventis Pharmaceuticals Inc., USA
SO PCT Int. Appl., 85 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2003006444	A2	20030123	WO 2002-US21525	20020709
	WO 2003006444	A3	20040311		
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	CA 2453169	AA	20030123	CA 2002-2453169	20020709
	US 2003087832	A1	20030508	US 2002-191718	20020709
	US 6777577	B2	20040817		
	EP 1423373	A2	20040602	EP 2002-749842	20020709
	EP 1423373	B1	20051019		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, SK				
	JP 2005504023	T2	20050210	JP 2003-512216	20020709
	AT 307125	E	20051115	AT 2002-749842	20020709
	ES 2247357	T3	20060301	ES 2002-2749842	20020709
	US 2004204582	A1	20041014	US 2004-835630	20040430
PRAI	US 2001-304020P	P	20010709		
	GB 2001-27615	A	20011119		
	US 2002-191718	A3	20020709		
	WO 2002-US21525	W	20020709		

OS MARPAT 138:122861

L11 ANSWER 18 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2002:946561 CAPLUS
DN 138:24739
TI Benzodiazepine bradykinin antagonists
IN Wood, Michael R.; Bock, Mark G.; Su, Dai-Shi; Kuduk, Scott D.; Han, Wei; Dorsey, Bruce D.
PA Merck & Co., Inc., USA
SO PCT Int. Appl., 71 pp.
CODEN: PIXXD2
DT Patent
LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2002099388	A2	20021212	WO 2002-US21065	20020603
	WO 2002099388	A3	20030501		

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

PRAI US 2001-296644P P 20010607
OS MARPAT 138:24739

L11 ANSWER 19 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2002:736230 CAPLUS
DN 137:263060
TI Preparation of heterocyclic compounds as αvβ3 integrin inhibitors
IN Morie, Toshiya; Iwama, Seiji; Notake, Mitsue; Kitano, Tomoko
PA Dainippon Pharmaceutical Co., Ltd., Japan
SO PCT Int. Appl., 115 pp.
CODEN: PIXXD2
DT Patent
LA Japanese
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2002074743	A1	20020926	WO 2002-JP2391	20020314

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

	EP 1371646	A1	20031217	EP 2002-705159	20020314

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR

	US 2004106622	A1	20040603	US 2003-472236	20030922
PRAI	JP 2001-79029	A	20010319		
	WO 2002-JP2391	W	20020314		

OS MARPAT 137:263060
RE.CNT 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 20 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2002:736222 CAPLUS
DN 137:262953
TI Preparation of biurethanes as inhibitors of blood-coagulation factor Xa and VIIa
IN Mederski, Werner; Cezanne, Bertram; Dorsch, Dieter; Tsaklakidis, Christos; Gleitz, Johannes; Barnes, Christopher
PA Merck Patent G.m.b.H., Germany
SO PCT Int. Appl., 33 pp.
CODEN: PIXXD2

DT Patent
LA German
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2002074735	A2	20020926	WO 2002-EP2095	20020227
	WO 2002074735	A3	20031127		
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	DE 10113402	A1	20020926	DE 2001-10113402	20010320
	CA 2441427	AA	20020926	CA 2002-2441427	20020227
	HU 200303512	A2	20040128	HU 2003-3512	20020227
	EP 1385818	A2	20040204	EP 2002-722151	20020227
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
	CN 1498206	A	20040519	CN 2002-806758	20020227
	JP 2004531494	T2	20041014	JP 2002-573744	20020227
	US 2004097550	A1	20040520	US 2003-472084	20030917
	US 6943179	B2	20050913		
	ZA 2003008060	A	20040723	ZA 2003-8060	20031016
PRAI	DE 2001-10113402	A	20010320		
	WO 2002-EP2095	W	20020227		

OS MARPAT 137:262953

L11 ANSWER 21 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2000:824211 CAPLUS
DN 134:4764
TI Preparation of 3-(benzoylamino)propionic acid derivatives as glucagon antagonists/inverse agonists
IN Ling, Anthony; Plewe, Michael Bruno; Truesdale, Larry Kenneth; Lau, Jesper; Madsen, Peter; Sams, Christian; Behrens, Carsten; Vagner, Josef; Christensen, Inge Thoger; Lundt, Behrend Frederik; Sidelmann, Ulla Grove; Thogersen, Henning
PA Novo Nordisk A/S, Den.; Agouron Pharmaceuticals, Inc.
SO PCT Int. Appl., 564 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2000069810	A1	20001123	WO 2000-DK264	20000516
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW				
	RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
	US 6503949	B1	20000516	US 2000-572553	20000516
	CA 2373892	AA	20001123	CA 2000-2373892	20000516
	EP 1183229	A1	20020306	EP 2000-926725	20000516
	EP 1183229	B1	20051026		

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO

	BR 2000010651	A	20020319	BR 2000-10651	20000516
	HU 200201033	A2	20020729	HU 2002-1033	20000516
	JP 2002544254	T2	20021224	JP 2000-618228	20000516
	AT 307798	E	20051115	AT 2000-926725	20000516
	ES 2250128	T3	20060416	ES 2000-926725	20000516
	ZA 2001008560	A	20020613	ZA 2001-8560	20011018
	NO 2001005607	A	20020117	NO 2001-5607	20011116
	US 2003220350	A1	20031127	US 2002-233851	20020830
	US 6875760	B2	20050405		
	US 2005203108	A1	20050915	US 2004-980199	20041103
PRAI	DK 1999-684	A	19990517		
	DK 2000-478	A	20000321		
	US 1999-134415P	P	19990517		
	US 2000-191685P	P	20000323		
	US 2000-572553	A3	20000516		
	WO 2000-DK264	W	20000516		
	US 2002-233851	A3	20020830		

OS MARPAT 134:4764
RE.CNT 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 22 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2000:756524 CAPLUS
DN 133:321878
TI Preparation of cyclic protein tyrosine kinase inhibitors
IN Das, Jagabandhu; Padmanabha, Ramesh; Chen, Ping; Norris, Derek J.; Doweyko, Arthur M. P.; Barrish, Joel C.; Wityak, John
PA Bristol-Myers Squibb Co., USA
SO PCT Int. Appl., 300 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2000062778	A1	20001026	WO 2000-US9753	20000412
	W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZA, ZW				
	RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
	CA 2366932	AA	20001026	CA 2000-2366932	20000412
	AU 2000042338	A5	20001102	AU 2000-42338	20000412
	AU 779089	B2	20050106		
	EP 1169038	A1	20020109	EP 2000-922102	20000412
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
	BR 2000009721	A	20020213	BR 2000-9721	20000412
	TR 200102969	T2	20020821	TR 2001-2969	20000412
	JP 2002542193	T2	20021210	JP 2000-611914	20000412
	HU 200202708	A2	20021228	HU 2002-2708	20000412
	NZ 513639	A	20040227	NZ 2000-513639	20000412
	RU 2260592	C2	20050920	RU 2001-130452	20000412
	ZA 2001007204	A	20021202	ZA 2001-7204	20010830
	NO 2001004970	A	20011210	NO 2001-4970	20011012
	US 2005261305	A1	20051124	US 2005-138793	20050525
	US 2005288303	A1	20051229	US 2005-138942	20050526

	US 2006079563	A1	20060413	US 2005-271626	20051110
PRAI	US 1999-129510P	P	19990415		
	WO 2000-US9753	W	20000412		
	US 2000-548929	A1	20000413		
	US 2003-378373	A1	20030303		

OS MARPAT 133:321878
RE.CNT 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 23 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2000:401817 CAPLUS
DN 133:30667
TI Heteroaryl-containing thiourea derivatives useful as inhibitors of herpes viruses
IN Bloom, Jonathan David; Digrandi, Martin Joseph; Dushin, Russell George; Lang, Stanley Albert; O'Hara, Bryan Mark
PA American Home Products Corporation, USA
SO PCT Int. Appl., 164 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 3

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2000034269	A1	20000615	WO 1999-US28892	19991206
	W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW				
	RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
	US 6166028	A	20001226	US 1999-444782	19991122
	US 6197803	B1	20010306	US 1999-447006	19991122
	US 6201013	B1	20010313	US 1999-444075	19991122
	CA 2351390	AA	20000615	CA 1999-2351390	19991206
	EP 1140913	A1	20011010	EP 1999-965143	19991206
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
	TR 200101598	T2	20011022	TR 2001-200101598	19991206
	BR 9916042	A	20011204	BR 1999-16042	19991206
	HU 200104758	A2	20020429	HU 2001-4758	19991206
	JP 2002531558	T2	20020924	JP 2000-586716	19991206
	AU 756043	B2	20030102	AU 2000-31122	19991206
	US 6262082	B1	20010717	US 2000-669483	20000925
	US 6271236	B1	20010807	US 2000-669943	20000926
	ZA 2001004373	A	20020918	ZA 2001-4373	20010528
	NO 2001002836	A	20010808	NO 2001-2836	20010608
	BG 105580	A	20020131	BG 2001-105580	20010608
	US 2003036653	A1	20030220	US 2002-99695	20020315
	US 6555561	B2	20030429		
PRAI	US 1998-208540	A	19981209		
	US 1998-150692P	P	19981209		
	US 1998-150698P	P	19981209		
	US 1998-155192P	P	19981209		
	US 1998-155240P	P	19981209		
	US 1998-208164	A	19981209		
	US 1998-208561	A	19981209		
	US 1999-444782	A3	19991122		
	WO 1999-US28892	W	19991206		
	US 2000-669535	A3	20000926		

OS MARPAT 133:30667
RE.CNT 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 24 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2000:401816 CAPLUS
DN 133:30666
TI Aryl- and heteroaryl-substituted thiourea derivatives useful as inhibitors of herpes viruses
IN Bloom, Jonathan David; Digrandi, Martin Joseph; Dushin, Russell George; Lang, Stanley Albert; O'Hara, Bryan Mark
PA American Home Products Corporation, USA
SO PCT Int. Appl., 159 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2000034268	A1	20000615	WO 1999-US28838	19991206
	W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW				
	RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
	CA 2350996	AA	20000615	CA 1999-2350996	19991206
	BR 9915993	A	20010904	BR 1999-15993	19991206
	EP 1137647	A1	20011004	EP 1999-965131	19991206
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
	HU 200104611	A2	20020429	HU 2001-4611	19991206
	JP 2002531557	T2	20020924	JP 2000-586715	19991206
	ZA 2001004318	A	20020826	ZA 2001-4318	20010525
	NO 2001002837	A	20010719	NO 2001-2837	20010608
PRAI	US 1998-207961	A	19981209		
	WO 1999-US28838	W	19991206		

OS MARPAT 133:30666
RE.CNT 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 25 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2000:401809 CAPLUS
DN 133:30657
TI Heterocyclic carboxamide-containing thiourea derivatives containing a substituted phenylenediamine group, useful as inhibitors of herpes viruses
IN Bloom, Jonathan David; Curran, Kevin Joseph; Digrandi, Martin Joseph; Dushin, Russell George; Jones, Thomas Richard; Lang, Stanley Albert; Ross, Adma Antonia; Terefenko, Eugene Anthony; O'Hara, Bryan Mark
PA American Home Products Corporation, USA
SO PCT Int. Appl., 159 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 3

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2000034261	A2	20000615	WO 1999-US28916	19991206
	WO 2000034261	A3	20020131		
	W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU,				

CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL,
IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA,
MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI,
SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW
RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE,
DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF,
CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

	US 6166028	A	20001226	US 1999-444782	19991122
	US 6197803	B1	20010306	US 1999-447006	19991122
	US 6201013	B1	20010313	US 1999-444075	19991122
	CA 2351690	AA	20000615	CA 1999-2351690	19991206
	EP 1144399	A2	20011017	EP 1999-967213	19991206
	EP 1144399	A3	20020911		

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
IE, SI, LT, LV, FI, RO

	BR 9916043	A	20011204	BR 1999-16043	19991206
	JP 2002533301	T2	20021008	JP 2000-586708	19991206
	HU 200203405	A2	20030228	HU 2002-3405	19991206
	US 6262082	B1	20010717	US 2000-669483	20000925
	US 6271236	B1	20010807	US 2000-669943	20000926
	ZA 2001004322	A	20021025	ZA 2001-4322	20010525
	NO 2001002835	A	20010719	NO 2001-2835	20010608
	US 2003036653	A1	20030220	US 2002-99695	20020315
	US 6555561	B2	20030429		
PRAI	US 1998-208164	A	19981209		
	US 1998-150692P	P	19981209		
	US 1998-150698P	P	19981209		
	US 1998-155192P	P	19981209		
	US 1998-155240P	P	19981209		
	US 1998-208540	A	19981209		
	US 1998-208561	A	19981209		
	US 1999-444782	A3	19991122		
	WO 1999-US28916	W	19991206		
	US 2000-669535	A3	20000926		

OS MARPAT 133:30657

L11 ANSWER 26 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2000:401808 CAPLUS
DN 133:30588
TI Alpha-methylbenzyl-containing thiourea derivatives containing a phenylenediamine group, useful as inhibitors of herpes viruses
IN Bloom, Jonathan David; Curran, Kevin Joseph; Digrandi, Martin Joseph; Dushin, Russell George; Lang, Stanley Albert; Norton, Emily Boucher; Ross, Adma Antonia; O'Hara, Bryan Mark
PA American Home Products Corporation, USA
SO PCT Int. Appl., 168 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2000034260	A2	20000615	WO 1999-US28839	19991206
	WO 2000034260	A3	20000908		

W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU,
CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL,
IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA,
MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI,
SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW
RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE,
DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF,
CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

	CA 2350833	AA	20000615	CA 1999-2350833	19991206
	BR 9916084	A	20010904	BR 1999-16084	19991206
	EP 1137645	A2	20011004	EP 1999-963022	19991206
	EP 1137645	B1	20040526		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
	HU 200104492	A2	20020429	HU 2001-4492	19991206
	JP 2002531555	T2	20020924	JP 2000-586707	19991206
	NZ 512135	A	20031219	NZ 1999-512135	19991206
	AT 267824	E	20040615	AT 1999-963022	19991206
	PT 1137645	T	20040930	PT 1999-963022	19991206
	ES 2221470	T3	20041216	ES 1999-963022	19991206
	ZA 2001004376	A	20020828	ZA 2001-4376	20010528
	NO 2001002833	A	20010802	NO 2001-2833	20010608
PRAI	US 1998-208902	A	19981209		
	WO 1999-US28839	W	19991206		

OS MARPAT 133:30588

L11 ANSWER 27 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2000:401806 CAPLUS
DN 133:30733
TI Heterocyclic carboxamide-containing thiourea derivatives containing a phenylenediamine group, useful as inhibitors of herpes viruses
IN Bloom, Jonathan David; Curran, Kevin Joseph; Digrandi, Martin Joseph; Dushin, Russell George; Jones, Thomas Richard; Lang, Stanley Albert; Ross, Adma Antonia; Terefenko, Eugene Anthony; O'Hara, Bryan Mark
PA American Home Products Corporation, USA
SO PCT Int. Appl., 188 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2000034258	A2	20000615	WO 1999-US28842	19991206
	WO 2000034258	A3	20011129		
	W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW				
	RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
	CA 2350767	AA	20000615	CA 1999-2350767	19991206
	BR 9916046	A	20011002	BR 1999-16046	19991206
	EP 1144397	A2	20011017	EP 1999-963023	19991206
	EP 1144397	A3	20020911		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
	TR 200101664	T2	20020321	TR 2001-200101664	19991206
	HU 200200232	A2	20020529	HU 2002-232	19991206
	JP 2002531554	T2	20020924	JP 2000-586705	19991206
	ZA 2001004377	A	20021220	ZA 2001-4377	20010528
	NO 2001002832	A	20010807	NO 2001-2832	20010608
	BG 105581	A	20011231	BG 2001-105581	20010608
PRAI	US 1998-208559	A	19981209		
	WO 1999-US28842	W	19991206		

OS MARPAT 133:30733

L11 ANSWER 28 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2000:401786 CAPLUS

DN 133:30587

TI Benzamide-containing aryl thiourea derivatives useful as inhibitors of herpes viruses

IN Bloom, Jonathan David; Curran, Kevin Joseph; Digrandi, Martin Joseph; Dushin, Russell George; Lang, Stanley Albert; Norton, Emily Boucher; Ross, Adma Antonia; O'Hara, Bryan Mark

PA American Home Products Corporation, USA

SO PCT Int. Appl., 169 pp.
CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 3

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2000034238	A1	20000615	WO 1999-US28837	19991206
	W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW				
	RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
	US 6166028	A	20001226	US 1999-444782	19991122
	US 6197803	B1	20010306	US 1999-447006	19991122
	US 6201013	B1	20010313	US 1999-444075	19991122
	US 6207715	B1	20010327	US 1999-444897	19991122
	US 6255349	B1	20010703	US 1999-444734	19991122
	US 6262090	B1	20010717	US 1999-444896	19991122
	US 6335350	B1	20020101	US 1999-447005	19991122
	CA 2351403	AA	20000615	CA 1999-2351403	19991206
	BR 9916086	A	20010904	BR 1999-16086	19991206
	EP 1137632	A1	20011004	EP 1999-963021	19991206
	EP 1137632	B1	20040728		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
	TR 200101597	T2	20011022	TR 2001-200101597	19991206
	HU 200104763	A2	20020429	HU 2001-4763	19991206
	JP 2002531545	T2	20020924	JP 2000-586686	19991206
	NZ 512108	A	20030926	NZ 1999-512108	19991206
	AT 272052	E	20040815	AT 1999-963021	19991206
	PT 1137632	T	20041130	PT 1999-963021	19991206
	ES 2224733	T3	20050301	ES 1999-963021	19991206
	US 6262082	B1	20010717	US 2000-669483	20000925
	US 6271236	B1	20010807	US 2000-669943	20000926
	US 6403617	B1	20020611	US 2000-669535	20000926
	US 6407123	B1	20020618	US 2000-670180	20000926
	US 6426355	B1	20020730	US 2000-671486	20000927
	US 6407249	B1	20020618	US 2000-684011	20001229
	US 6410571	B1	20020625	US 2000-684773	20001229
	US 2002026055	A1	20020228	US 2001-804510	20010312
	US 6380243	B2	20020430		
	US 2001039348	A1	20011108	US 2001-845428	20010430
	US 6462055	B2	20021008		
	ZA 2001004144	A	20020821	ZA 2001-4144	20010521
	NO 2001002838	A	20010808	NO 2001-2838	20010608
	BG 105583	A	20011231	BG 2001-105583	20010608
	US 2003036653	A1	20030220	US 2002-99695	20020315
	US 6555561	B2	20030429		
PRAI	US 1998-208561	A	19981209		
	US 1998-228805P	P	19981209		
	US 1998-228808P	P	19981209		

	US 1998-228809P	P	19981209
	US 1998-150692P	P	19981209
	US 1998-150698P	P	19981209
	US 1998-155192P	P	19981209
	US 1998-155240P	P	19981209
	US 1998-208164	A	19981209
	US 1998-208540	A	19981209
	US 1999-444734	A3	19991122
	US 1999-444782	A3	19991122
	US 1999-444896	A3	19991122
	US 1999-447006	A3	19991122
	WO 1999-US28837	W	19991206
	US 2000-669535	A3	20000926

OS MARPAT 133:30587
RE.CNT 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 29 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2000:401785 CAPLUS
DN 133:30586
TI Acetamide and substituted acetamide-containing aryl thiourea derivatives useful as inhibitors of herpes viruses
IN Bloom, Jonathan David; Digrandi, Martin Joseph; Dushin, Russell George; Lang, Stanley Albert; O'Hara, Bryan Mark
PA American Home Products Corporation, USA
SO PCT Int. Appl., 159 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2000034237	A2	20000615	WO 1999-US28844	19991206
	WO 2000034237	A3	20001123		
	W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW				
	RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
	CA 2350899	AA	20000615	CA 1999-2350899	19991206
	EP 1137633	A2	20011004	EP 1999-965132	19991206
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
	BR 9916041	A	20011204	BR 1999-16041	19991206
	HU 200104944	A2	20020429	HU 2001-4944	19991206
	JP 2002531544	T2	20020924	JP 2000-586685	19991206
	ZA 2001004142	A	20021025	ZA 2001-4142	20010521
	NO 2001002834	A	20010807	NO 2001-2834	20010608
PRAI	US 1998-208316	A	19981209		
	WO 1999-US28844	W	19991206		

OS MARPAT 133:30586

L11 ANSWER 30 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 1999:9703 CAPLUS
DN 130:81404
TI Piperidinylazacycloalkylmethylureas as α1A adrenergic receptor antagonists
IN Patane, Michael A.; Bock, Mark G.
PA Merck & Co., Inc., USA

SO PCT Int. Appl., 143 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 9857640	A1	19981223	WO 1998-US12672	19980618
W: AL, AM, AU, AZ, BA, BB, BG, BR, BY, CA, CN, CU, CZ, EE, GE, GW, HU, ID, IL, IS, JP, KG, KR, KZ, LC, LK, LR, LT, LV, MD, MG, MK, MN, MX, NO, NZ, PL, RO, RU, SG, SI, SK, SL, TJ, TM, TR, TT, UA, US, UZ, VN, YU, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG				
EP 1019052	A1	20000719	EP 1998-931353	19980617
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, PT, IE, FI				
US 6143750	A	20001107	US 1998-98780	19980617
CA 2294346	AA	19981223	CA 1998-2294346	19980618
AU 9881501	A1	19990104	AU 1998-81501	19980618
JP 2002508764	T2	20020319	JP 1999-504780	19980618
PRAI US 1997-50960P	P	19970618		
GB 1998-231	A	19980106		
WO 1998-US12672	W	19980618		

OS MARPAT 130:81404
RE.CNT 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 31 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 1999:9701 CAPLUS
DN 130:81519
TI Preparation of [2-(piperidin-4-yl)aminoethylcarbamoyl] substituted 1,2,3,4-tetrahydropyrimidines and oxazolidines as alpha 1a adrenergic receptor antagonists
IN Patane, Michael A.; Bock, Mark G.; Newton, Randall C.
PA Merck & Co., Inc., USA
SO PCT Int. Appl., 175 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 9857638	A1	19981223	WO 1998-US12567	19980617
W: AL, AM, AU, AZ, BA, BB, BG, BR, BY, CA, CN, CU, CZ, EE, GE, GW, HU, ID, IL, IS, JP, KG, KR, KZ, LC, LK, LR, LT, LV, MD, MG, MK, MN, MX, NO, NZ, PL, RO, RU, SG, SI, SK, SL, TJ, TM, TR, TT, UA, US, UZ, VN, YU, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG				
CA 2294590	AA	19981223	CA 1998-2294590	19980617
AU 9879726	A1	19990104	AU 1998-79726	19980617
EP 1023068	A1	20000802	EP 1998-930307	19980617
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, PT, IE, FI				
JP 2002511085	T2	20020409	JP 1999-504715	19980617
US 6376503	B1	20020423	US 1998-97947	19980617
PRAI US 1997-50959P	P	19970618		
GB 1998-217	A	19980107		
WO 1998-US12567	W	19980617		

OS MARPAT 130:81519
RE.CNT 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD

ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 32 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 1997:357099 CAPLUS
DN 127:81761
TI Cyclic homopentapeptides. 1. Analogs of tuberactinomycins and capreomycin with activity against vancomycin-resistant enterococci and Pasteurella
AU Dirlam, J. P.; Belton, A. M.; Birsner, N. C.; Brooks, R. R.; Chang, S.-P.; Chandrasekaran, R. Y.; Clancy, J.; Cronin, B. J.; Dirlam, B. P.; Finegan, S. M.; Froshauer, S. A.; Girard, A. E.; Hayashi, S. F.; Howe, R. J.; Kane, J. C.; Kamicker, B. J.; Kaufman, S. A.; Kolosko, N. L.; LeMay, M. A.; Linde, R. G., II; Lyssikatos, J. P.; MacLelland, C. P.; Magee, T. V.; Massa, M. A.; Miller, S. A.; Minich, M. L.; Perry, D. A.; Petitpas, J. W.; Reese, C. P.; Seibel, S. B.; Su, W.-G.; Sweeney, K. T.; Whipple, D. A.; Yang, B. V.
CS Central Research Division, Pfizer Inc., Groton, CT, 06340, USA
SO Bioorganic & Medicinal Chemistry Letters (1997), 7(9), 1139-1144
CODEN: BMCLE8; ISSN: 0960-894X
PB Elsevier
DT Journal
LA English
OS CASREACT 127:81761
RE.CNT 12 THERE ARE 12 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 33 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 1996:509383 CAPLUS
DN 125:167546
TI Preparation of aniline derivatives as nitrogen monoxide synthase inhibitors
IN Honda, Toshio; Makino, Toshihiko; Nagafuji, Toshiaki; Kitoh, Yasushi; Kimura, Nobuaki
PA Chugai Seiyaku Kabushiki Kaisha, Japan
SO PCT Int. Appl., 384 pp.
CODEN: PIXXD2
DT Patent
LA Japanese
FAN.CNT 2

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 9618608	A1	19960620	WO 1995-JP2540	19951212
W: AL, AM, AU, BB, BG, BR, BY, CA, CN, CZ, EE, FI, GE, HU, IS, KE, KG, KR, KZ, LK, LR, LS, LT, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, RO, RU, SD, SG, SI, SK, TJ, TM, TT, UA, UG, US, UZ, VN				
RW: KE, LS, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG				
CA 2206005	AA	19960620	CA 1995-2206005	19951212
CA 2206005	C	20060502		
AU 9641240	A1	19960703	AU 1996-41240	19951212
AU 705152	B2	19990513		
EP 798292	A1	19971001	EP 1995-939418	19951212
EP 798292	B1	20041103		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE				
BR 9510006	A	19971111	BR 1995-10006	19951212
NZ 296594	A	20000228	NZ 1995-296594	19951212
RU 2167858	C2	20010527	RU 1997-111792	19951212
PL 183619	B1	20020628	PL 1995-320829	19951212
AT 281430	E	20041115	AT 1995-939418	19951212
TW 474909	B	20020201	TW 1995-84113596	19951219
US 6534546	B1	20030318	US 1997-849400	19970606
FI 9702460	A	19970811	FI 1997-2460	19970610

	NO 9702666	A	19970812	NO 1997-2666	19970610
	NO 310615	B1	20010730		
	LT 4343	B	19980525	LT 1997-119	19970710
	HK 1008867	A1	20020705	HK 1998-109613	19980801
PRAI	JP 1994-336795	A	19941212		
	JP 1995-113695	A	19950414		
	WO 1995-JP1135	A	19950607		
	WO 1995-JP2540	W	19951212		

OS MARPAT 125:167546

L11 ANSWER 34 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 1995:758682 CAPLUS
DN 123:169279
TI Preparation of cyclohexanediurea derivatives as ACAT inhibitors
IN Yamada, Toshihiro; Nobuhara, Yoichi; Takagi, Ichinari; Furumoto, Shiho; Kobayashi, Kazuhiro; Ikemoto, Kiyohito
PA Nissin Food Products Co., Ltd., Japan
SO PCT Int. Appl., 146 pp.
CODEN: PIXXD2
DT Patent
LA Japanese
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 9507258	A1	19950316	WO 1994-JP1475	19940907
W: AU, CA, KR, US				
RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
JP 07082232	A2	19950328	JP 1993-226247	19930910
JP 3286745	B2	20020527		
CA 2171295	AA	19950316	CA 1994-2171295	19940907
AU 9476235	A1	19950327	AU 1994-76235	19940907
AU 680941	B2	19970814		
EP 718281	A1	19960626	EP 1994-926366	19940907
EP 718281	B1	20010627		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, MC, NL, PT, SE				
AT 202555	E	20010715	AT 1994-926366	19940907
US 5733931	A	19980331	US 1996-617828	19960308
PRAI JP 1993-226247	A	19930910		
WO 1994-JP1475	W	19940907		

OS MARPAT 123:169279

L11 ANSWER 35 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 1993:179931 CAPLUS
DN 118:179931
TI Silver halide photographic material
IN Kato, Kazunobu
PA Fuji Photo Film Co., Ltd., Japan
SO Jpn. Kokai Tokkyo Koho, 36 pp.
CODEN: JKXXAF
DT Patent
LA Japanese
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI JP 04330432	A2	19921118	JP 1991-128214	19910502
US 5262274	A	19931116	US 1992-876386	19920430
PRAI JP 1991-128214	A	19910502		

L11 ANSWER 36 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 1992:214907 CAPLUS
DN 116:214907
TI Preparation of N-acetyl-N-phenylglycinanides as drugs

IN Bourzat, Jean Dominique; Capet, Marc; Cotrel, Claude; Guyon, Claude; Manfre, Franco; Roussel, Gerard
PA Rhone-Poulenc Rorer SA, Fr.
SO PCT Int. Appl., 145 pp.
CODEN: PIXXD2
DT Patent
LA French
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9112264	A1	19910822	WO 1991-FR87	19910206
	W: AU, CA, HU, JP, KR, NO, SU, US				
	RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LU, NL, SE				
	FR 2658196	A1	19910816	FR 1990-1553	19900209
	FR 2658196	B1	19920424		
	FR 2667319	A2	19920403	FR 1990-11916	19900927
	FR 2667319	B2	19921120		
	FR 2667863	A2	19920417	FR 1990-12594	19901012
	FR 2667863	B2	19921127		
	CA 2072981	AA	19910810	CA 1991-2072981	19910206
	AU 9173295	A1	19910903	AU 1991-73295	19910206
	AU 639081	B2	19930715		
	EP 514442	A1	19921125	EP 1991-903956	19910206
	EP 514442	B1	19940427		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE				
	HU 61575	A2	19930128	HU 1992-2585	19910206
	JP 05506643	T2	19930930	JP 1991-504069	19910206
	AT 104989	E	19940515	AT 1991-903956	19910206
	ES 2052372	T3	19940701	ES 1991-903956	19910206
	ZA 9100946	A	19911127	ZA 1991-946	19910208
	US 5382590	A	19950117	US 1992-867690	19920708
	NO 9203079	A	19920805	NO 1992-3079	19920805
PRAI	FR 1990-1553	A	19900209		
	FR 1990-11916	A	19900927		
	FR 1990-12594	A	19901012		
	EP 1991-903956	A	19910206		
	WO 1991-FR87	A	19910206		

OS MARPAT 116:214907

L11 ANSWER 37 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 1992:151348 CAPLUS
DN 116:151348
TI Preparation of aromatic diurea derivatives and their salts
IN Ito, Tokuki; Matsuda, Mitsuaki; Izumi, Yuichi
PA Yamanouchi Pharmaceutical Co., Ltd., Japan
SO Jpn. Kokai Tokkyo Koho, 5 pp.
CODEN: JKXXAF
DT Patent
LA Japanese
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 03255061	A2	19911113	JP 1990-50074	19900301
PRAI	JP 1990-50074		19900301		

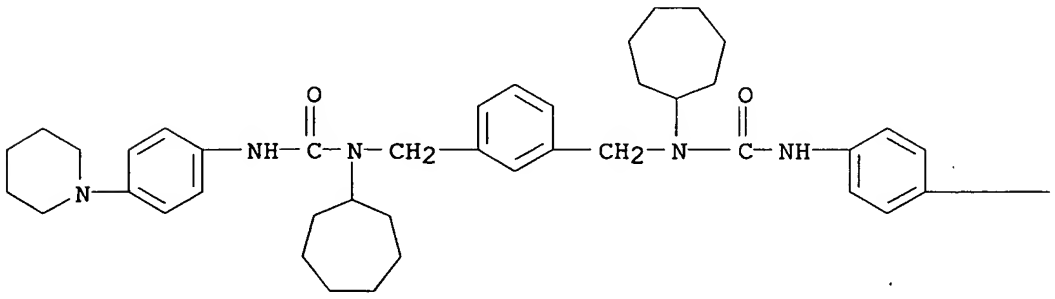
OS MARPAT 116:151348

=> d hitstr 37

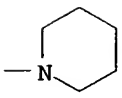
L11 ANSWER 37 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 139649-87-9P
RL: SPN (Synthetic preparation); PREP (Preparation)

(preparation of, as arteriosclerosis inhibitor)
RN 139649-87-9 CAPLUS
CN Urea, N,N''-[1,3-phenylenebis(methylene)]bis[N-cycloheptyl-N'-[4-(1-piperidinyl)phenyl]- (9CI) (CA INDEX NAME)

PAGE 1-A

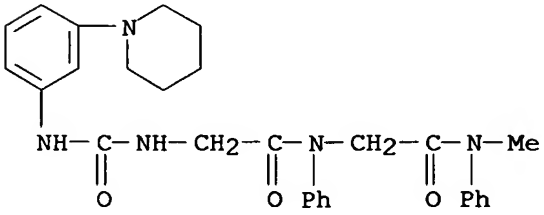


PAGE 1-B



=> d hitstr 36

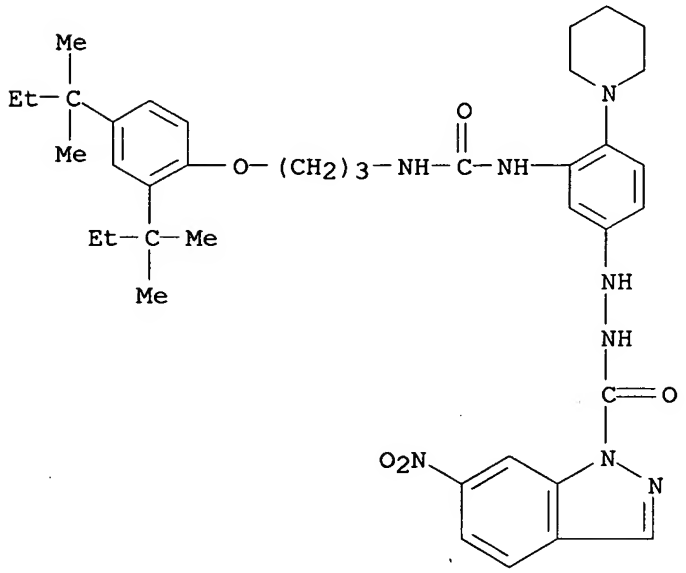
L11 ANSWER 36 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 138562-07-9P
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of, as antagonist of CCK and gastrin)
RN 138562-07-9 CAPLUS
CN Glycinamide, N-[[[3-(1-piperidinyl)phenyl]amino]carbonyl]glycyl-N-methyl-N,N2-diphenyl- (9CI) (CA INDEX NAME)



=> d hitstr 35

L11 ANSWER 35 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 146657-34-3
RL: TEM (Technical or engineered material use); USES (Uses)

(silver halide photog. materials containing)
RN 146657-34-3 CAPLUS
CN 1H-Indazole-1-carboxylic acid, 6-nitro-, 2-[3-[[[[3-[2,4-bis(1,1-dimethylpropyl)phenoxy]propyl]amino]carbonyl]amino]-4-(1-piperidinyl)phenyl]hydrazide (9CI) (CA INDEX NAME)

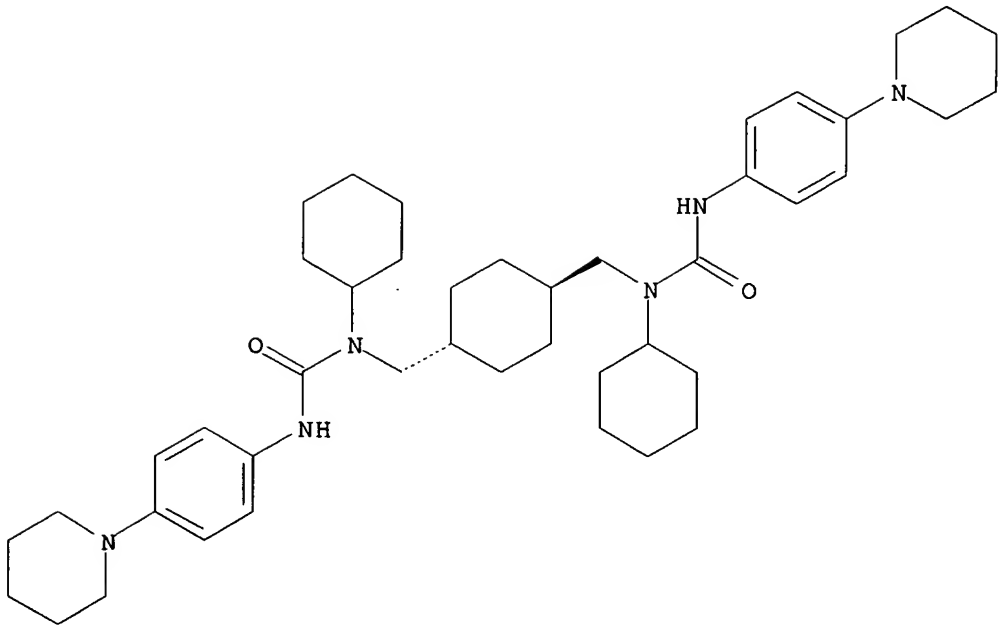


=> d hitstr 34

L11 ANSWER 34 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 166967-89-1P 166968-04-3P 166968-21-4P
166968-72-5P 166968-91-8P 166968-95-2P
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of cyclohexanediurea derivs. as ACAT inhibitors)
RN 166967-89-1 CAPLUS
CN Urea, N,N''-[1,4-cyclohexanediylbis(methylene)]bis[N-cyclohexyl-N'-[4-(1-piperidinyl)phenyl]-, dihydrochloride, trans- (9CI) (CA INDEX NAME)

Relative stereochemistry.

PAGE 1-A



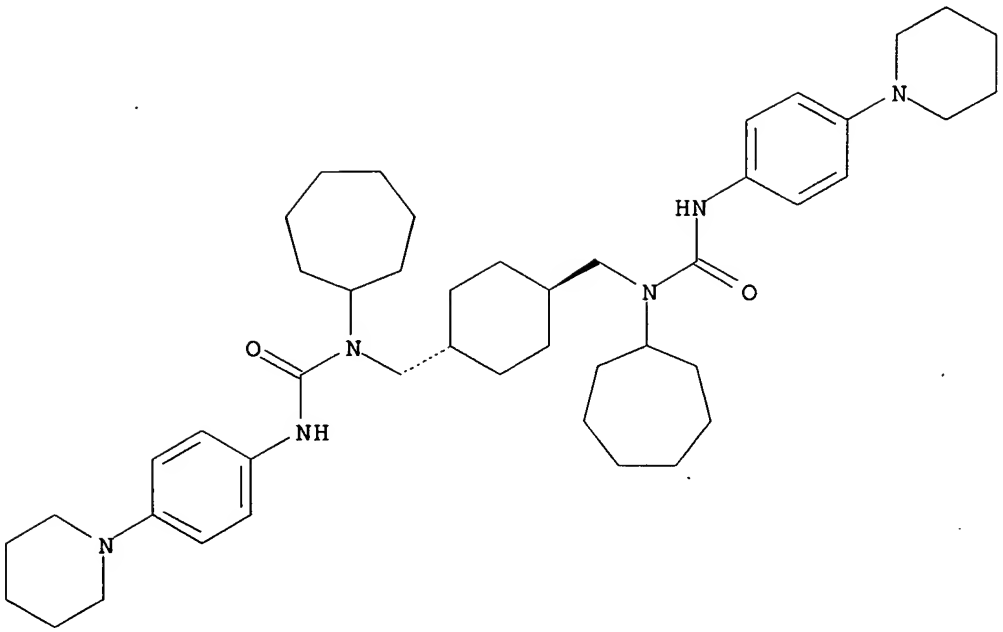
PAGE 2-A

●2 HCl

RN 166968-04-3 CAPLUS
CN Urea, N,N''-[1,4-cyclohexanediylbis(methylene)]bis[N-cycloheptyl-N'-[4-(1-piperidinyl)phenyl]-, dihydrochloride, trans- (9CI) (CA INDEX NAME)

Relative stereochemistry.

PAGE 1-A

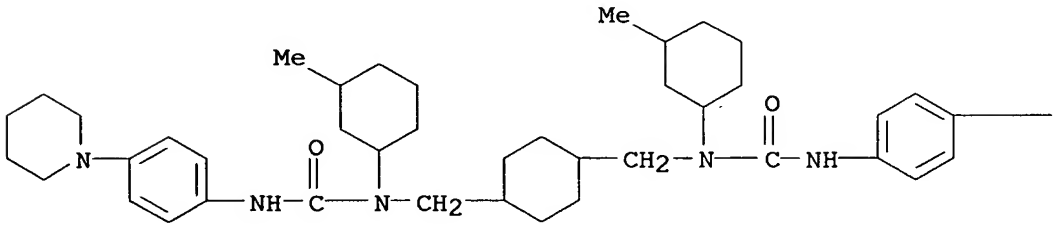


PAGE 2-A

●2 HCl

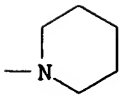
RN 166968-21-4 CAPLUS
CN Urea, N,N''-[1,4-cyclohexanediylbis(methylene)]bis[N-(3-methylcyclohexyl)-N'-[4-(1-piperidinyl)phenyl]-, dihydrochloride (9CI) (CA INDEX NAME)

PAGE 1-A



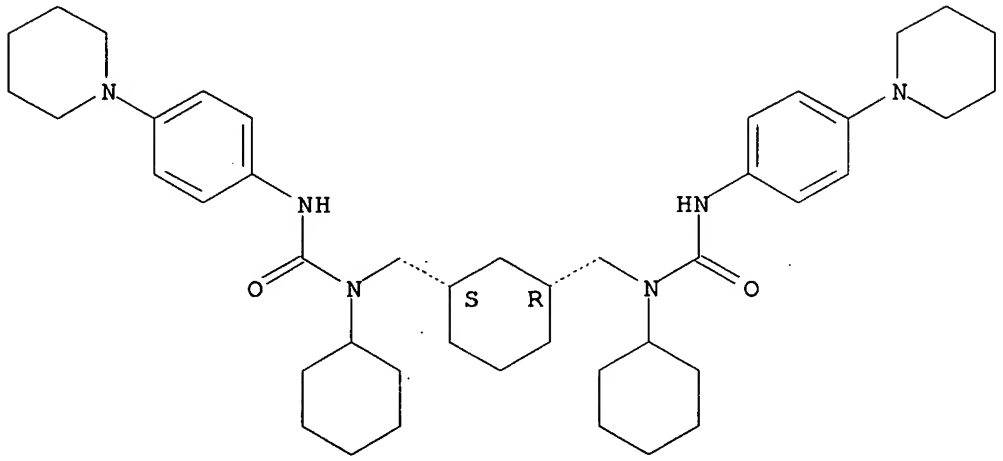
●2 HCl

PAGE 1-B



RN 166968-72-5 CAPLUS
CN Urea, N,N''-[1,3-cyclohexanediylbis(methylene)]bis[N-cyclohexyl-N'-[4-(1-piperidinyl)phenyl]-, dihydrochloride, cis- (9CI) (CA INDEX NAME)

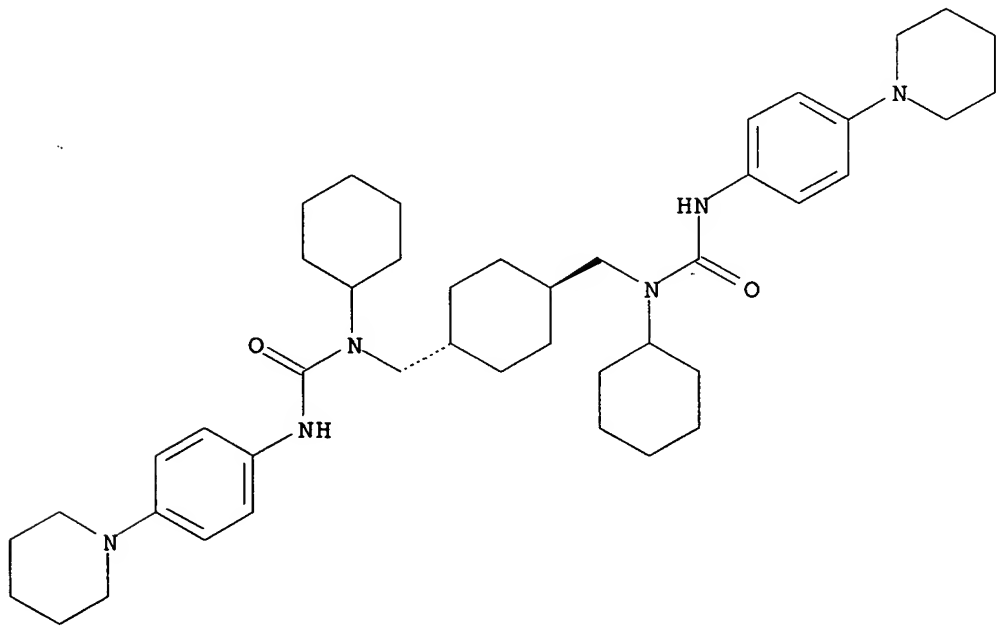
Relative stereochemistry.



●2 HCl

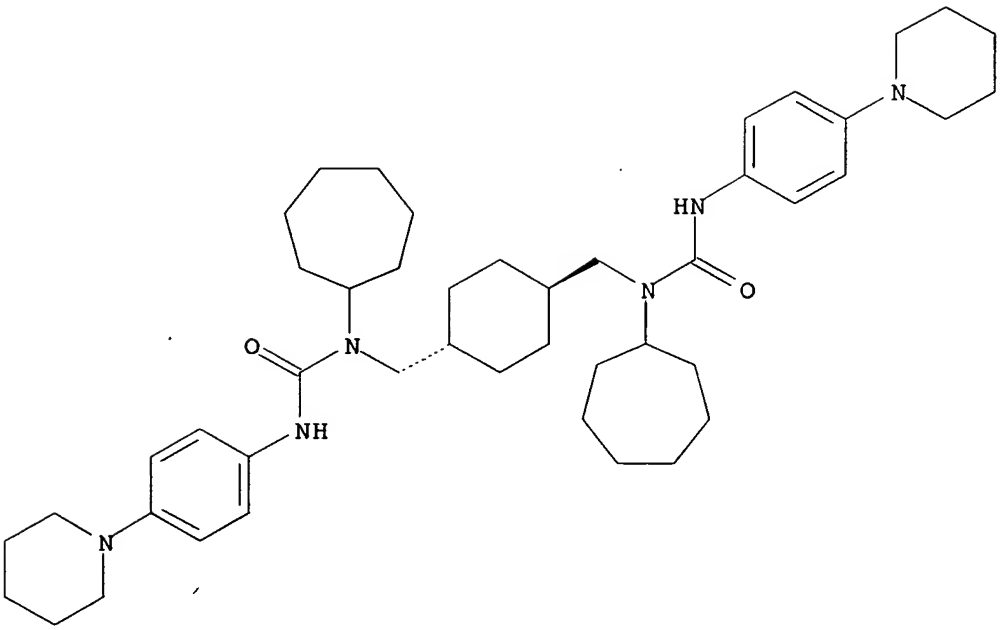
RN 166968-91-8 CAPLUS
CN Urea, N,N''-[1,4-cyclohexanediylbis(methylene)]bis[N-cyclohexyl-N'-[4-(1-piperidinyl)phenyl]-, trans- (9CI) (CA INDEX NAME)

Relative stereochemistry.



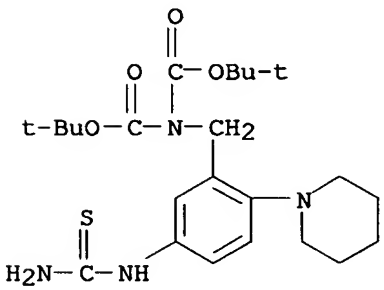
RN 166968-95-2 CAPLUS
CN Urea, N,N''-[1,4-cyclohexanediylbis(methylene)]bis[N-cycloheptyl-N'-[4-(1-piperidinyl)phenyl]-, trans- (9CI) (CA INDEX NAME)

Relative stereochemistry.

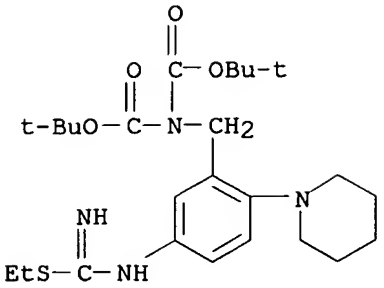


=> d hitstr 33

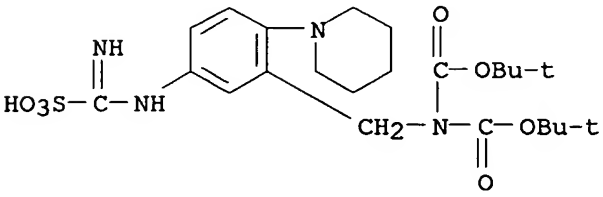
L11 ANSWER 33 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 180149-65-9P 180149-66-0P 180149-68-2P
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of aniline derivs. as nitrogen monoxide synthase inhibitors)
RN 180149-65-9 CAPLUS
CN Imidodicarbonic acid, [[5-[(aminothioxomethyl)amino]-2-(1-piperidinyl)phenyl]methyl]-, bis(1,1-dimethylethyl) ester (9CI) (CA INDEX NAME)



RN 180149-66-0 CAPLUS
CN Imidodicarbonic acid, [[5-[[(ethylthio)iminomethyl]amino]-2-(1-piperidinyl)phenyl]methyl]-, bis(1,1-dimethylethyl) ester (9CI) (CA INDEX NAME)



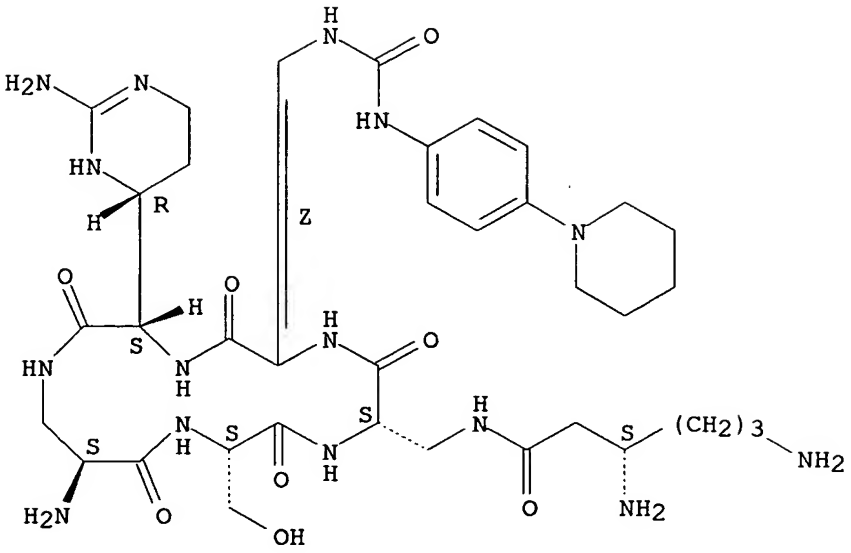
RN 180149-68-2 CAPLUS
CN Methanesulfonic acid, [[3-[[bis[(1,1-dimethylethoxy)carbonyl]amino]methyl]-4-(1-piperidinyl)phenyl]amino]imino- (9CI) (CA INDEX NAME)



=> d hitstr 32

L11 ANSWER 32 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 191670-61-8P
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation)
(preparation of tuberactinomycin and capreomycin analogs with antibacterial activity against vancomycin-resistant enterococci and Pasteurella)
RN 191670-61-8 CAPLUS
CN Cyclo[3-[[(3S)-3,6-diamino-1-oxohexyl]amino]-L-alanyl-(2Z)-2,3-didehydro-3-[[[[4-(1-piperidinyl)phenyl]amino]carbonyl]amino]alanyl-(2S)-2-[(4R)-2-amino-1,4,5,6-tetrahydro-4-pyrimidinyl]glycyl-(2S)-2-amino-β-alanyl-L-seryl] (9CI) (CA INDEX NAME)

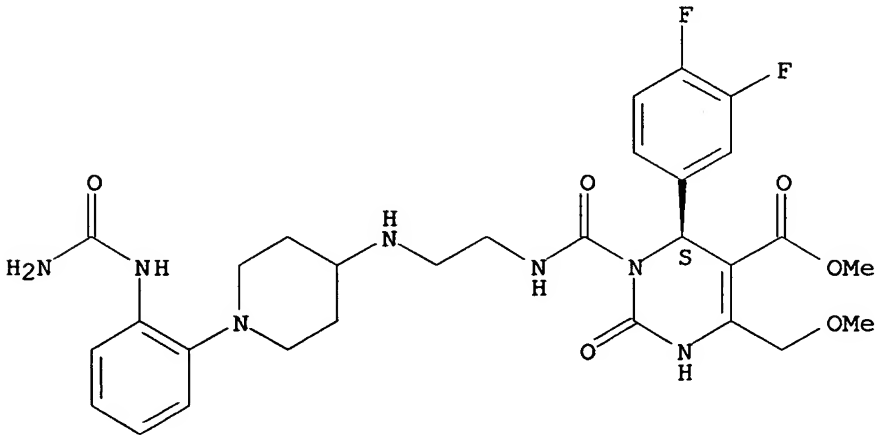
Absolute stereochemistry.
Double bond geometry as shown.



=> d hitstr 31

L11 ANSWER 31 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 218609-73-5P 218609-81-5P
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of [2-(piperidin-4-yl)aminoethylcarbamoyl] substituted 1,2,3,4-tetrahydropyrimidines and oxazolidines as alpha 1a adrenergic receptor antagonists)
RN 218609-73-5 CAPLUS
CN 5-Pyrimidinecarboxylic acid, 1-[[[2-[[1-[2-[(aminocarbonyl)amino]phenyl]-4-piperidinyl]amino]ethyl]amino]carbonyl]-6-(3,4-difluorophenyl)-1,2,3,6-tetrahydro-4-(methoxymethyl)-2-oxo-, methyl ester, (6S)- (9CI) (CA INDEX NAME)

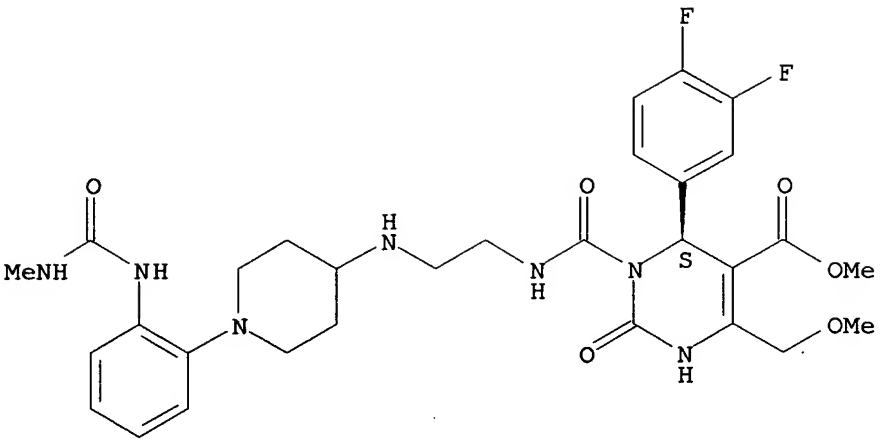
Absolute stereochemistry.



RN 218609-81-5 CAPLUS
CN 5-Pyrimidinecarboxylic acid, 6-(3,4-difluorophenyl)-1,2,3,6-tetrahydro-4-(methoxymethyl)-1-[[[2-[[1-[2-[[(methylamino)carbonyl]amino]phenyl]-4-piperidinyl]amino]ethyl]amino]carbonyl]-2-oxo-, methyl ester, (6S)- (9CI)

(CA INDEX NAME)

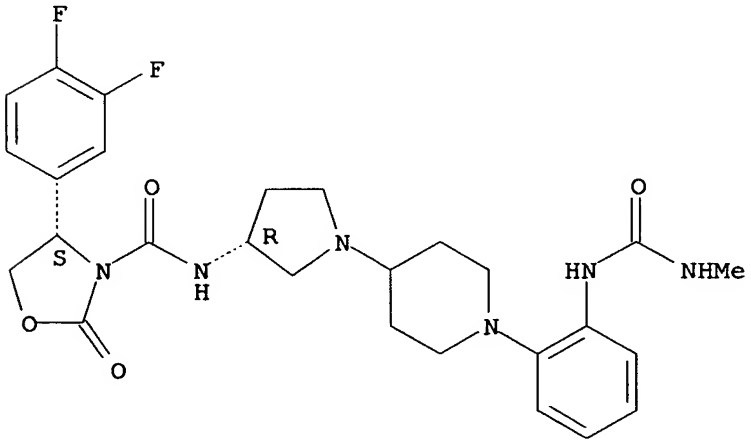
Absolute stereochemistry.



=> d hitstr 30

L11 ANSWER 30 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 218430-78-5P 218430-83-2P
RL: SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of piperidinylazacycloalkylmethylureas as α1A adrenergic receptor antagonists)
RN 218430-78-5 CAPLUS
CN 3-Oxazolidinecarboxamide, 4-(3,4-difluorophenyl)-N-[(3R)-1-[1-[2-[[(methylamino)carbonyl]amino]phenyl]-4-piperidinyl]-3-pyrrolidinyl]-2-oxo-, hydrochloride (5:11), (4S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



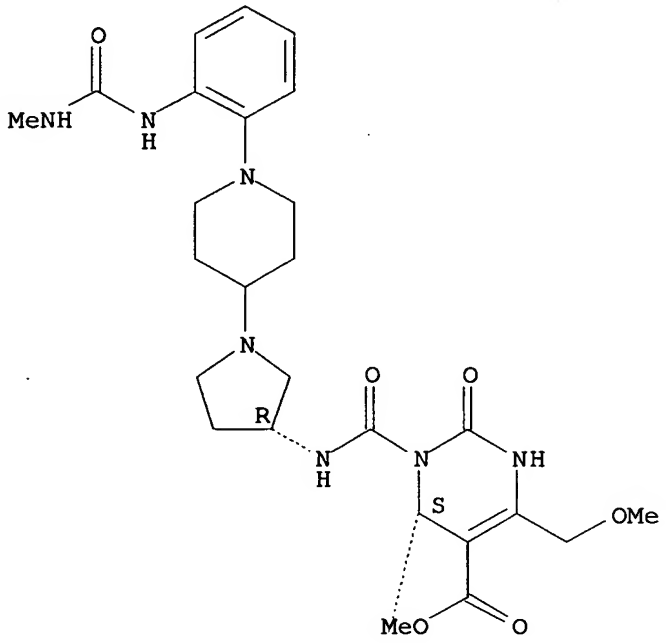
●11/5 HCl

RN 218430-83-2 CAPLUS
CN 5-Pyrimidinecarboxylic acid, 6-(3,4-difluorophenyl)-1,2,3,6-tetrahydro-4-

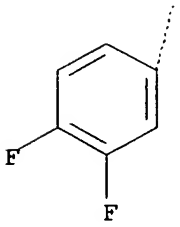
(methoxymethyl)-1-[[[(3R)-1-[1-[2-[[(methylamino)carbonyl]amino]phenyl]-4-piperidinyl]-3-pyrrolidinyl]amino]carbonyl]-2-oxo-, methyl ester, (6S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A

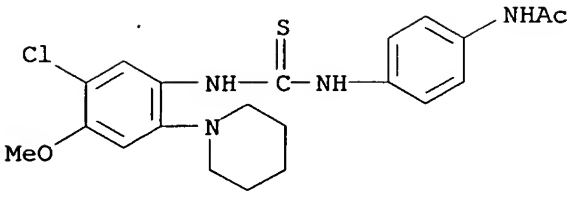


PAGE 2-A

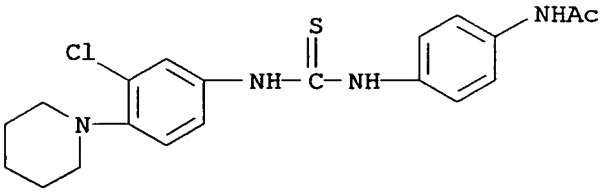


=> d hitstr 29

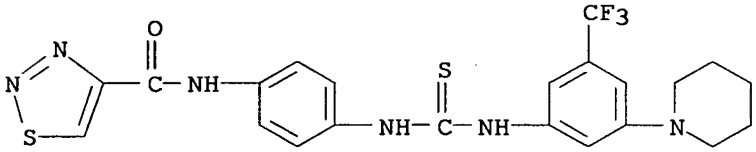
L11 ANSWER 29 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 273386-61-1P 273386-77-9P 273389-91-6P
273390-26-4P
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(target compound; preparation of acetamide-containing aryl thiourea derivs. as
inhibitors of herpes viruses)
RN 273386-61-1 CAPLUS
CN Acetamide, N-[4-[[[[5-chloro-4-methoxy-2-(1-piperidinyl)phenyl]amino]thioxomethyl]amino]phenyl]- (9CI) (CA INDEX NAME)



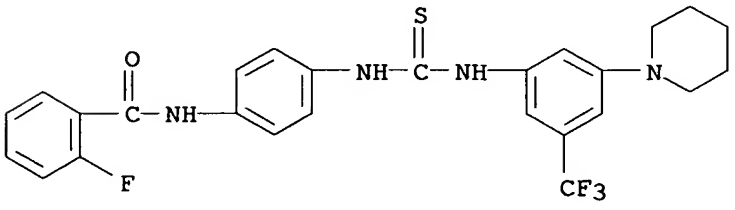
RN 273386-77-9 CAPLUS
CN Acetamide, N-[4-[[[[3-chloro-4-(1-piperidinyl)phenyl]amino]thioxomethyl]amino]phenyl]- (9CI) (CA INDEX NAME)



RN 273389-91-6 CAPLUS
CN 1,2,3-Thiadiazole-4-carboxamide, N-[4-[[[[3-(1-piperidinyl)-5-(trifluoromethyl)phenyl]amino]thioxomethyl]amino]phenyl]- (9CI) (CA INDEX NAME)



RN 273390-26-4 CAPLUS
CN Benzamide, 2-fluoro-N-[4-[[[[3-(1-piperidinyl)-5-(trifluoromethyl)phenyl]amino]thioxomethyl]amino]phenyl]- (9CI) (CA INDEX NAME)



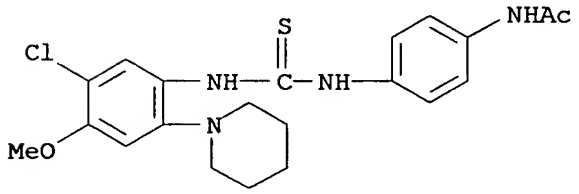
=> d hitstr 28

L11 ANSWER 28 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 273386-61-1P 273386-77-9P 273389-91-6P
273390-26-4P
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use);

BIOL (Biological study); PREP (Preparation); USES (Uses)
(target compound; preparation of benzamide-containing aryl thiourea derivs. as
inhibitors of herpes viruses)

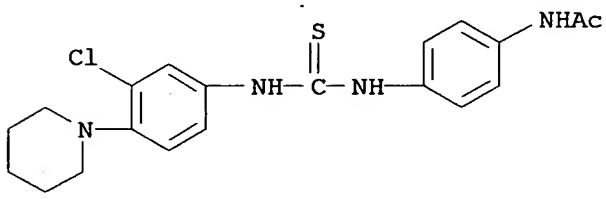
RN 273386-61-1 CAPLUS

CN Acetamide, N-[4-[[[[5-chloro-4-methoxy-2-(1-piperidinyl)phenyl]amino]thioxomethyl]amino]phenyl]- (9CI) (CA INDEX NAME)



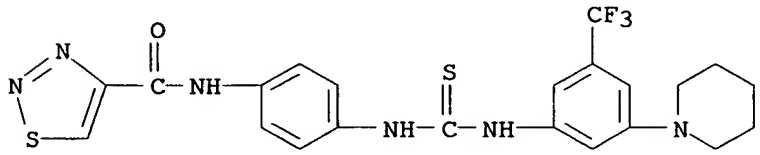
RN 273386-77-9 CAPLUS

CN Acetamide, N-[4-[[[[3-chloro-4-(1-piperidinyl)phenyl]amino]thioxomethyl]amino]phenyl]- (9CI) (CA INDEX NAME)



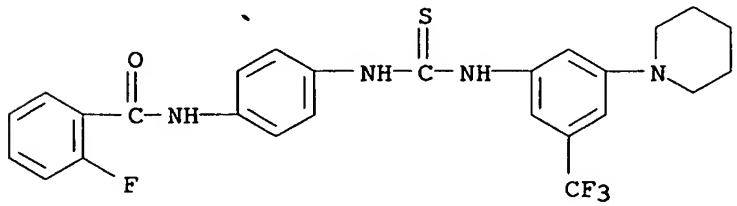
RN 273389-91-6 CAPLUS

CN 1,2,3-Thiadiazole-4-carboxamide, N-[4-[[[[3-(1-piperidinyl)-5-(trifluoromethyl)phenyl]amino]thioxomethyl]amino]phenyl]- (9CI) (CA INDEX NAME)



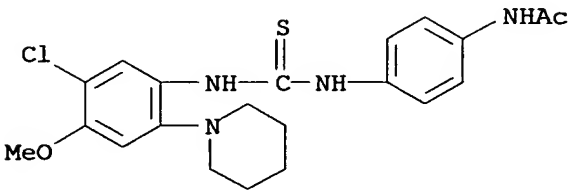
RN 273390-26-4 CAPLUS

CN Benzamide, 2-fluoro-N-[4-[[[[3-(1-piperidinyl)-5-(trifluoromethyl)phenyl]amino]thioxomethyl]amino]phenyl]- (9CI) (CA INDEX NAME)

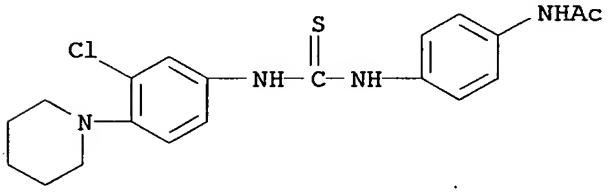


=> d hitstr 27

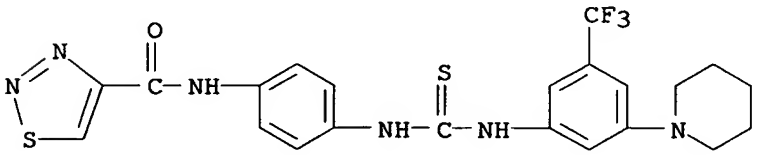
L11 ANSWER 27 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 273386-61-1P 273386-77-9P 273389-91-6P
273390-26-4P
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(target compound; preparation of heterocyclic carboxamide-containing thiourea
derivs. as inhibitors of herpes viruses)
RN 273386-61-1 CAPLUS
CN Acetamide, N-[4-[[[[5-chloro-4-methoxy-2-(1-piperidinyl)phenyl]amino]thioxomethyl]amino]phenyl]- (9CI) (CA INDEX NAME)



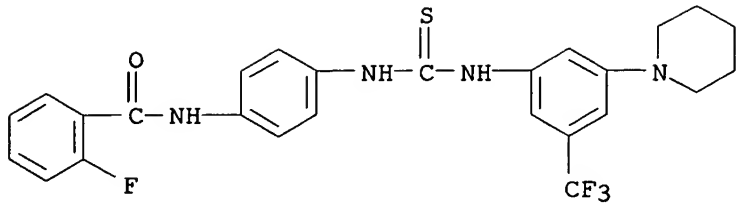
RN 273386-77-9 CAPLUS
CN Acetamide, N-[4-[[[[3-chloro-4-(1-piperidinyl)phenyl]amino]thioxomethyl]amino]phenyl]- (9CI) (CA INDEX NAME)



RN 273389-91-6 CAPLUS
CN 1,2,3-Thiadiazole-4-carboxamide, N-[4-[[[[3-(1-piperidinyl)-5-(trifluoromethyl)phenyl]amino]thioxomethyl]amino]phenyl]- (9CI) (CA INDEX NAME)

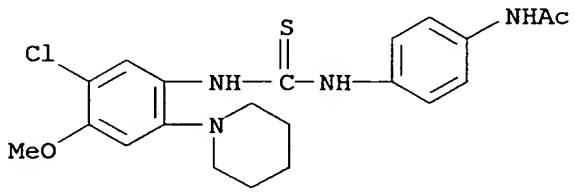


RN 273390-26-4 CAPLUS
CN Benzamide, 2-fluoro-N-[4-[[[[3-(1-piperidinyl)-5-(trifluoromethyl)phenyl]amino]thioxomethyl]amino]phenyl]- (9CI) (CA INDEX NAME)

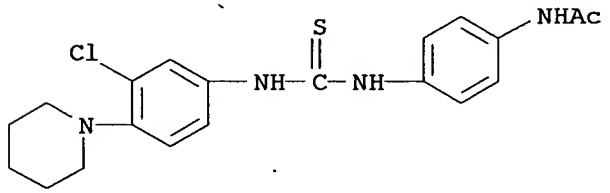


=> d hitstr 26

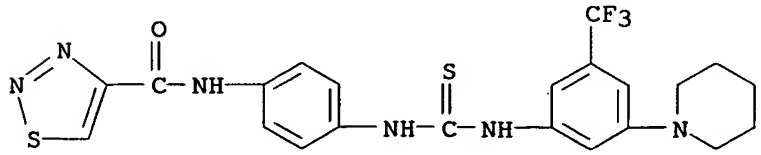
L11 ANSWER 26 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 273386-61-1P 273386-77-9P 273389-91-6P
273390-26-4P
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(target compound; preparation of α-methylbenzyl-containing thiourea derivs. as inhibitors of herpes viruses)
RN 273386-61-1 CAPLUS
CN Acetamide, N-[4-[[[[5-chloro-4-methoxy-2-(1-piperidinyl)phenyl]amino]thioxomethyl]amino]phenyl]- (9CI) (CA INDEX NAME)



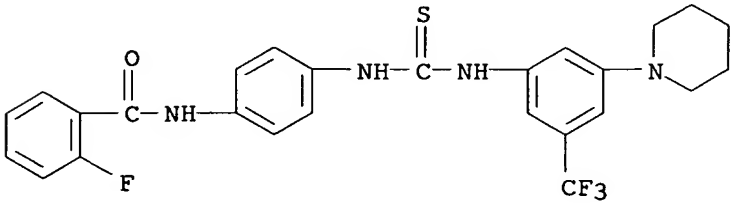
RN 273386-77-9 CAPLUS
CN Acetamide, N-[4-[[[[3-chloro-4-(1-piperidinyl)phenyl]amino]thioxomethyl]amino]phenyl]- (9CI) (CA INDEX NAME)



RN 273389-91-6 CAPLUS
CN 1,2,3-Thiadiazole-4-carboxamide, N-[4-[[[[3-(1-piperidinyl)-5-(trifluoromethyl)phenyl]amino]thioxomethyl]amino]phenyl]- (9CI) (CA INDEX NAME)

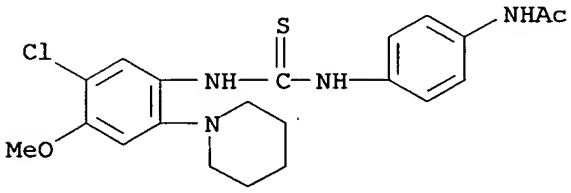


RN 273390-26-4 CAPLUS
CN Benzamide, 2-fluoro-N-[4-[[[[3-(1-piperidinyl)-5-(trifluoromethyl)phenyl]amino]thioxomethyl]amino]phenyl]- (9CI) (CA INDEX NAME)

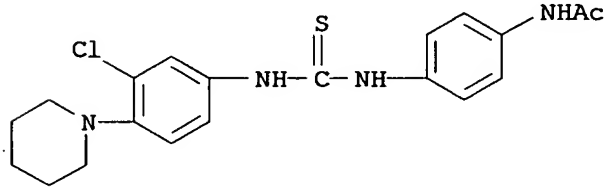


=> d hitstr 25

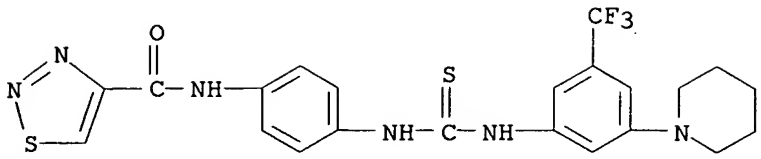
L11 ANSWER 25 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 273386-61-1P 273386-77-9P 273389-91-6P
273390-26-4P
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(target compound; preparation of heterocyclic carboxamide-containing and phenylenediamine-containing thiourea derivs. as inhibitors of herpes viruses)
RN 273386-61-1 CAPLUS
CN Acetamide, N-[4-[[[[5-chloro-4-methoxy-2-(1-piperidinyl)phenyl]amino]thioxomethyl]amino]phenyl]- (9CI) (CA INDEX NAME)



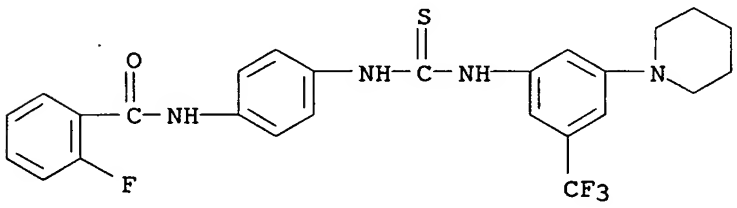
RN 273386-77-9 CAPLUS
CN Acetamide, N-[4-[[[[3-chloro-4-(1-piperidinyl)phenyl]amino]thioxomethyl]amino]phenyl]- (9CI) (CA INDEX NAME)



RN 273389-91-6 CAPLUS
CN 1,2,3-Thiadiazole-4-carboxamide, N-[4-[[[[3-(1-piperidinyl)-5-(trifluoromethyl)phenyl]amino]thioxomethyl]amino]phenyl]- (9CI) (CA INDEX NAME)

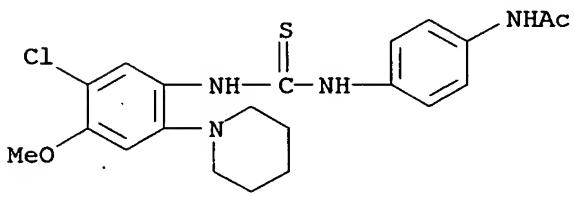


RN 273390-26-4 CAPLUS
CN Benzamide, 2-fluoro-N-[4-[[[[3-(1-piperidinyl)-5-(trifluoromethyl)phenyl]amino]thioxomethyl]amino]phenyl]- (9CI) (CA INDEX NAME)

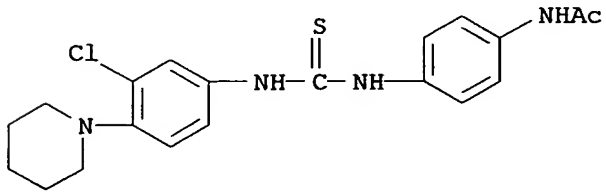


=> d hitstr 24

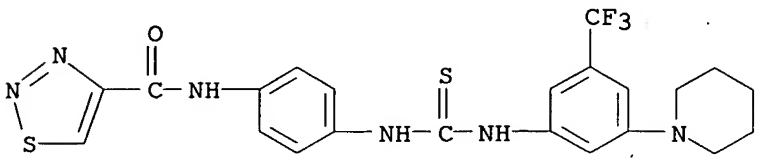
L11 ANSWER 24 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 273386-61-1P 273386-77-9P 273389-91-6P
273390-26-4P
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(target compound; preparation of heteroaryl thiourea derivs. as inhibitors of
herpes viruses)
RN 273386-61-1 CAPLUS
CN Acetamide, N-[4-[[[[5-chloro-4-methoxy-2-(1-piperidinyl)phenyl]amino]thioxomethyl]amino]phenyl]- (9CI) (CA INDEX NAME)



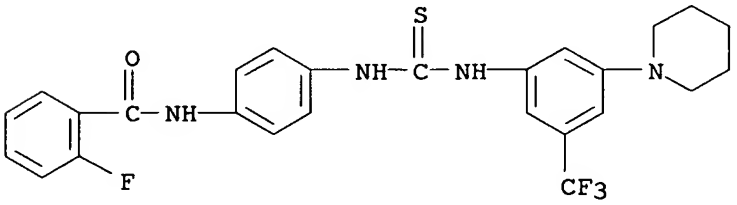
RN 273386-77-9 CAPLUS
CN Acetamide, N-[4-[[[[3-chloro-4-(1-piperidinyl)phenyl]amino]thioxomethyl]amino]phenyl]- (9CI) (CA INDEX NAME)



RN 273389-91-6 CAPLUS
CN 1,2,3-Thiadiazole-4-carboxamide, N-[4-[[[[3-(1-piperidinyl)-5-(trifluoromethyl)phenyl]amino]thioxomethyl]amino]phenyl]- (9CI) (CA INDEX NAME)

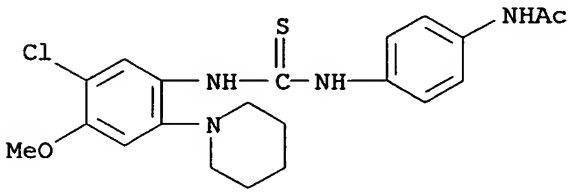


RN 273390-26-4 CAPLUS
CN Benzamide, 2-fluoro-N-[4-[[[[3-(1-piperidinyl)-5-(trifluoromethyl)phenyl]amino]thioxomethyl]amino]phenyl]- (9CI) (CA INDEX NAME)

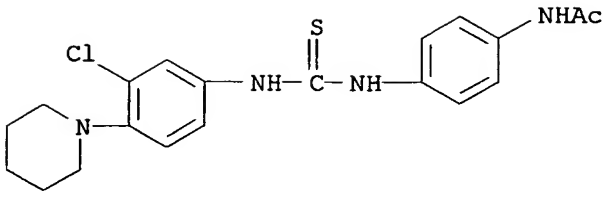


=> d hitstr 24

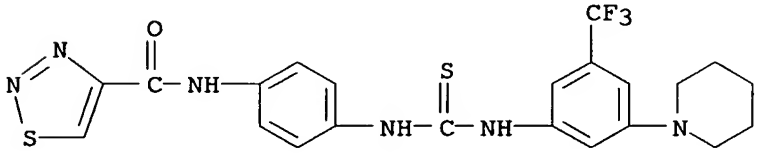
L11 ANSWER 24 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 273386-61-1P 273386-77-9P 273389-91-6P
273390-26-4P
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(target compound; preparation of heteroaryl thiourea derivs. as inhibitors of
herpes viruses)
RN 273386-61-1 CAPLUS
CN Acetamide, N-[4-[[[[5-chloro-4-methoxy-2-(1-piperidinyl)phenyl]amino]thioxomethyl]amino]phenyl]- (9CI) (CA INDEX NAME)



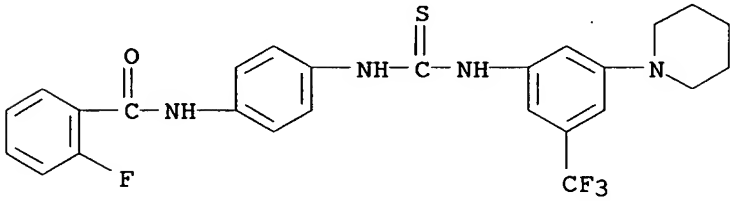
RN 273386-77-9 CAPLUS
CN Acetamide, N-[4-[[[[3-chloro-4-(1-piperidinyl)phenyl]amino]thioxomethyl]amino]phenyl]- (9CI) (CA INDEX NAME)



RN 273389-91-6 CAPLUS
CN 1,2,3-Thiadiazole-4-carboxamide, N-[4-[[[[3-(1-piperidinyl)-5-(trifluoromethyl)phenyl]amino]thioxomethyl]amino]phenyl]- (9CI) (CA INDEX NAME)

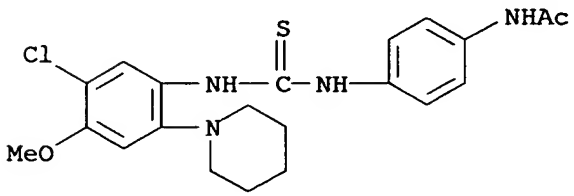


RN 273390-26-4 CAPLUS
CN Benzamide, 2-fluoro-N-[4-[[[[3-(1-piperidinyl)-5-(trifluoromethyl)phenyl]amino]thioxomethyl]amino]phenyl]- (9CI) (CA INDEX NAME)

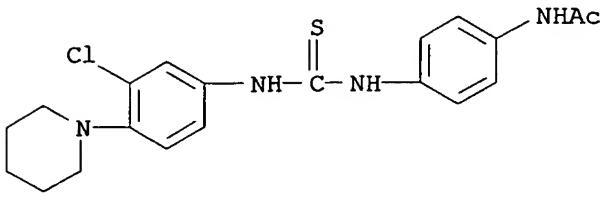


=> d hitstr 23

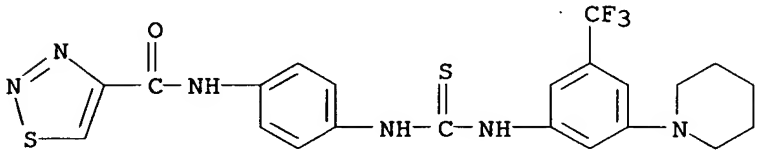
L11 ANSWER 23 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 273386-61-1P 273386-77-9P 273389-91-6P
273390-26-4P
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(target compound; preparation of heteroaryl-containing thiourea derivs. as inhibitors of herpes viruses)
RN 273386-61-1 CAPLUS
CN Acetamide, N-[4-[[[[5-chloro-4-methoxy-2-(1-piperidinyl)phenyl]amino]thioxomethyl]amino]phenyl]- (9CI) (CA INDEX NAME)



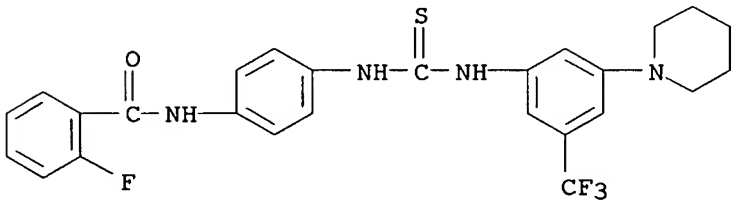
RN 273386-77-9 CAPLUS
CN Acetamide, N-[4-[[[[3-chloro-4-(1-piperidinyl)phenyl]amino]thioxomethyl]amino]phenyl]- (9CI) (CA INDEX NAME)



RN 273389-91-6 CAPLUS
CN 1,2,3-Thiadiazole-4-carboxamide, N-[4-[[[[3-(1-piperidinyl)-5-(trifluoromethyl)phenyl]amino]thioxomethyl]amino]phenyl]- (9CI) (CA INDEX NAME)

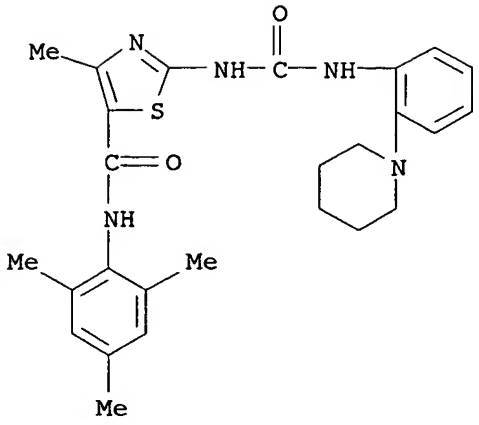


RN 273390-26-4 CAPLUS
CN Benzamide, 2-fluoro-N-[4-[[[[3-(1-piperidinyl)-5-(trifluoromethyl)phenyl]amino]thioxomethyl]amino]phenyl]- (9CI) (CA INDEX NAME)



=> d hitstr 22

L11 ANSWER 22 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 302960-70-9P
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of cyclic protein tyrosine kinase inhibitors)
RN 302960-70-9 CAPLUS
CN 5-Thiazolecarboxamide, 4-methyl-2-[[[[2-(1-piperidinyl)phenyl]amino]carbonyl]amino]-N-(2,4,6-trimethylphenyl)- (9CI) (CA INDEX NAME)



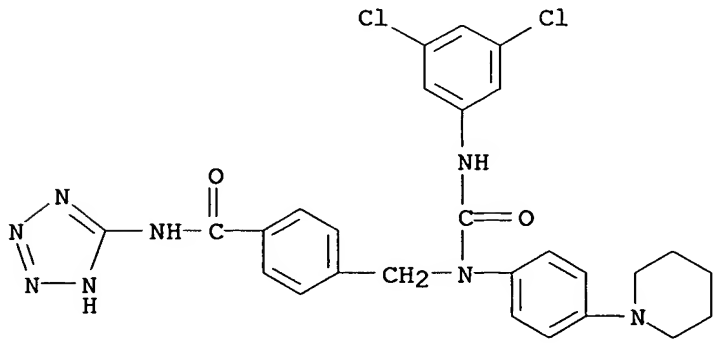
=> d hitstr 21

L11 ANSWER 21 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN

IT 307985-68-8P 307985-69-9P 307985-70-2P
307986-88-5P
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of 3-(benzoylamino)propionic acid derivs. as glucagon antagonists/inverse agonists)

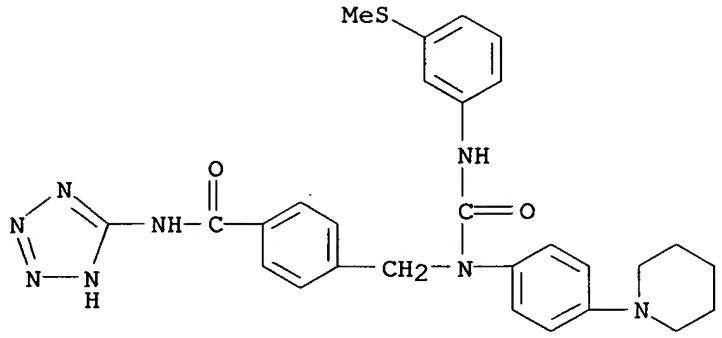
RN 307985-68-8 CAPLUS

CN Benzamide, 4-[[[[(3,5-dichlorophenyl)amino]carbonyl][4-(1-piperidinyl)phenyl]amino]methyl]-N-1H-tetrazol-5-yl- (9CI) (CA INDEX NAME)

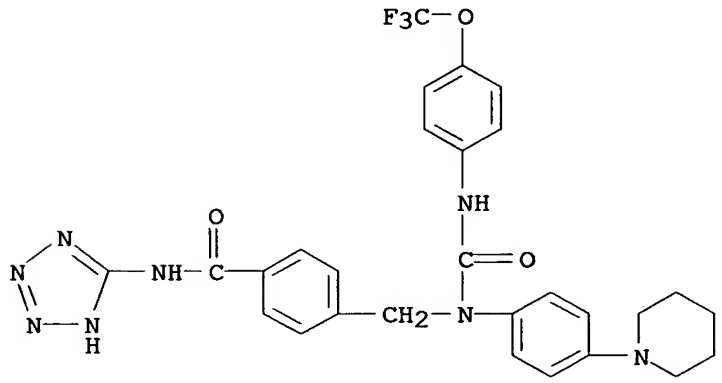


RN 307985-69-9 CAPLUS

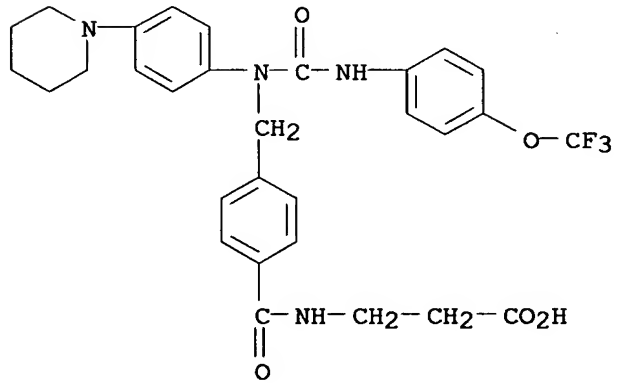
CN Benzamide, 4-[[[[[3-(methylthio)phenyl]amino]carbonyl][4-(1-piperidinyl)phenyl]amino]methyl]-N-1H-tetrazol-5-yl- (9CI) (CA INDEX NAME)



RN 307985-70-2 CAPLUS
CN Benzamide, 4-[[[4-(1-piperidinyl)phenyl][[[4-(trifluoromethoxy)phenyl]amino]carbonyl]amino]methyl]-N-1H-tetrazol-5-yl- (9CI) (CA INDEX NAME)



RN 307986-88-5 CAPLUS
CN β-Alanine, N-[4-[[[4-(1-piperidinyl)phenyl][[[4-(trifluoromethoxy)phenyl]amino]carbonyl]amino]methyl]benzoyl]- (9CI) (CA INDEX NAME)

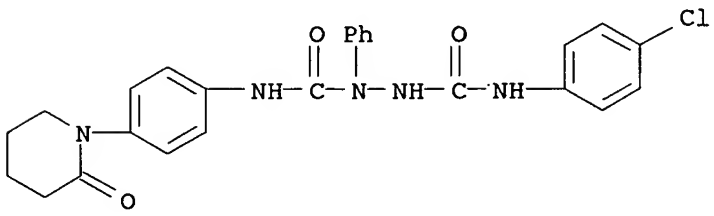


=> d hitstr 20

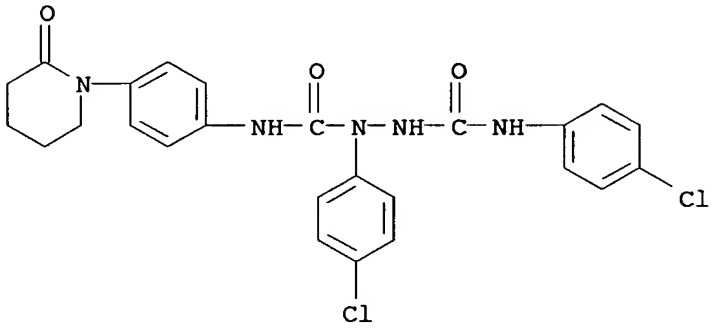
L11 ANSWER 20 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 461435-24-5P 461435-25-6P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of biurethanes as inhibitors of blood-coagulation factor Xa and VIIa)

RN 461435-24-5 CAPLUS
CN 1,2-Hydrazinedicarboxamide, N2-(4-chlorophenyl)-N1-[4-(2-oxo-1-piperidinyl)phenyl]-1-phenyl- (9CI) (CA INDEX NAME)

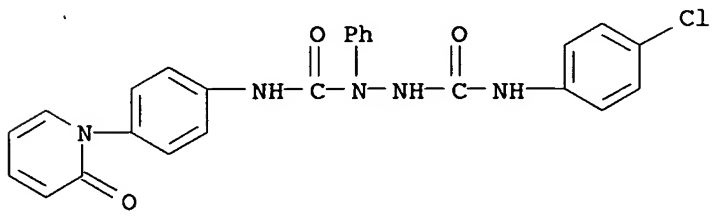


RN 461435-25-6 CAPLUS
CN 1,2-Hydrazinedicarboxamide, N2,1-bis(4-chlorophenyl)-N1-[4-(2-oxo-1-piperidinyl)phenyl]- (9CI) (CA INDEX NAME)

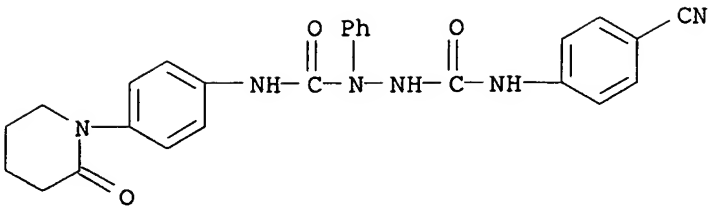


IT 461435-30-3 461435-34-7
RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
(preparation of biurethanes as inhibitors of blood-coagulation factor Xa and VIIa)

RN 461435-30-3 CAPLUS
CN 1,2-Hydrazinedicarboxamide, N2-(4-chlorophenyl)-N1-[4-(2-oxo-1(2H)-pyridinyl)phenyl]-1-phenyl- (9CI) (CA INDEX NAME)



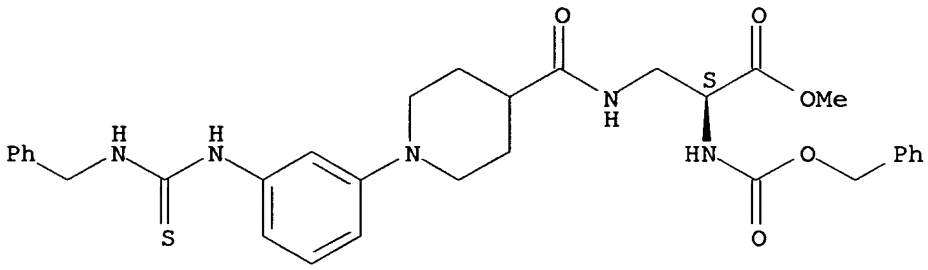
RN 461435-34-7 CAPLUS
CN 1,2-Hydrazinedicarboxamide, N2-(4-cyanophenyl)-N1-[4-(2-oxo-1-piperidinyl)phenyl]-1-phenyl- (9CI) (CA INDEX NAME)



=> d hitstr 19

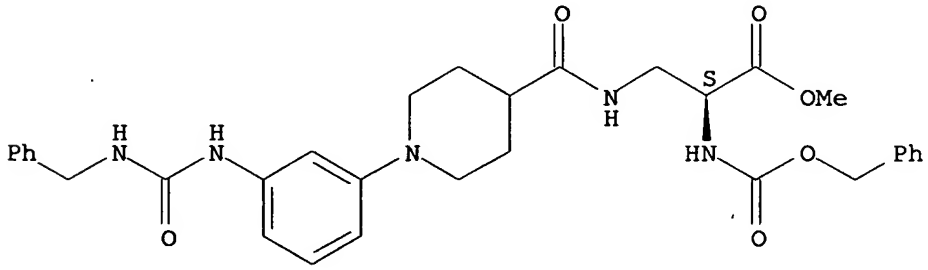
L11 ANSWER 19 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 461718-89-8P 461718-94-5P
RL: PAC (Pharmacological activity); RCT (Reactant); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); RACT (Reactant or reagent); USES (Uses)
(preparation of heterocyclic compds. as αvβ3 integrin inhibitors)
RN 461718-89-8 CAPLUS
CN L-Alanine, N-[(phenylmethoxy)carbonyl]-3-[[[1-[3-[[[(phenylmethyl)amino]thioxomethyl]amino]phenyl]-4-piperidinyl]carbonyl]amino]-, methyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.



RN 461718-94-5 CAPLUS
CN L-Alanine, N-[(phenylmethoxy)carbonyl]-3-[[[1-[3-[[[(phenylmethyl)amino]carbonyl]amino]phenyl]-4-piperidinyl]carbonyl]amino]-, methyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.



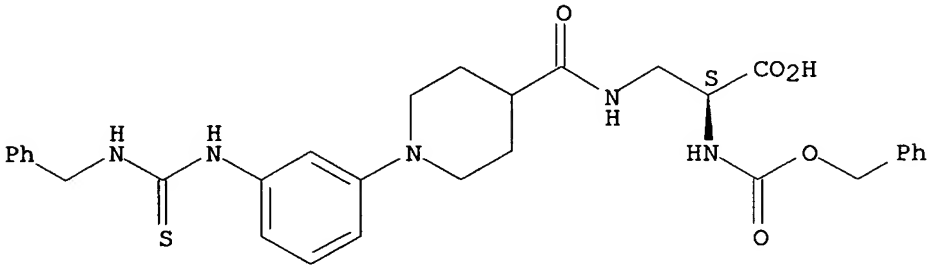
IT 461718-90-1P 461718-91-2P 461718-92-3P
461718-93-4P 461718-95-6P 461718-96-7P
461718-97-8P 461718-98-9P 461718-99-0P
461719-00-6P 461719-01-7P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU

(Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of heterocyclic compds. as αvβ3 integrin inhibitors)

RN 461718-90-1 CAPLUS

CN L-Alanine, N-[(phenylmethoxy)carbonyl]-3-[[[1-[3-[[[(phenylmethyl)amino]thioxomethyl]amino]phenyl]-4-piperidinyl]carbonyl]amino]- (9CI) (CA INDEX NAME)

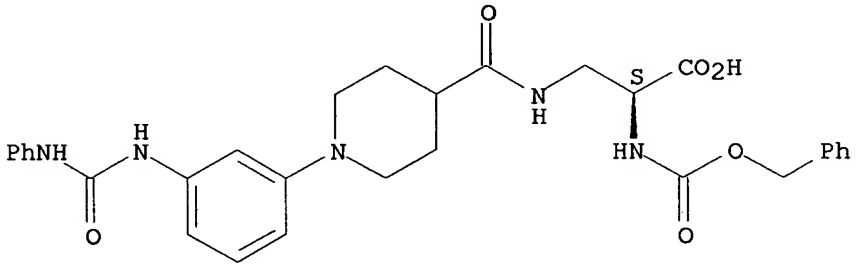
Absolute stereochemistry.



RN 461718-91-2 CAPLUS

CN L-Alanine, 3-[[[1-[3-[[(phenylamino)carbonyl]amino]phenyl]-4-piperidinyl]carbonyl]amino]-N-[(phenylmethoxy)carbonyl]- (9CI) (CA INDEX NAME)

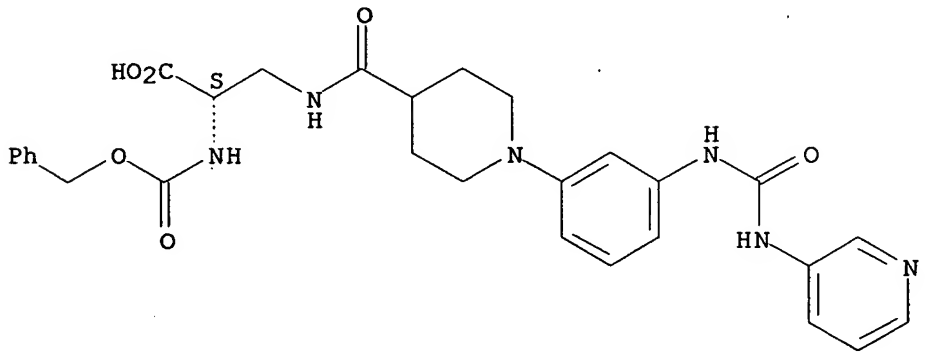
Absolute stereochemistry.



RN 461718-92-3 CAPLUS

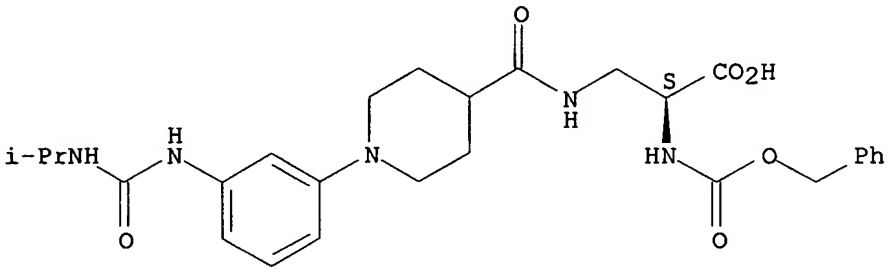
CN L-Alanine, N-[(phenylmethoxy)carbonyl]-3-[[[1-[3-[[(3-pyridinylamino)carbonyl]amino]phenyl]-4-piperidinyl]carbonyl]amino]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



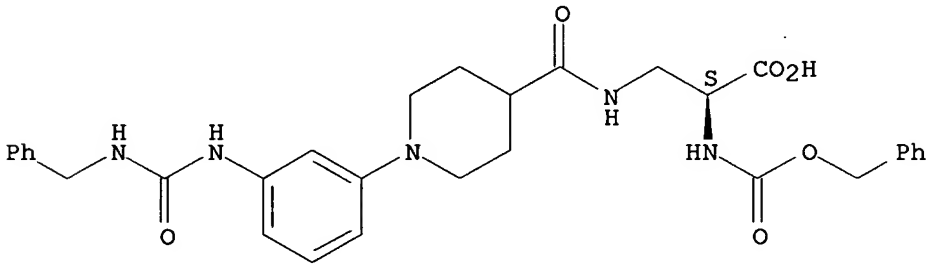
RN 461718-93-4 CAPLUS
CN L-Alanine, 3-[[[1-[3-[[[(1-methylethyl)amino]carbonyl]amino]phenyl]-4-piperidinyl]carbonyl]amino]-N-[(phenylmethoxy)carbonyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

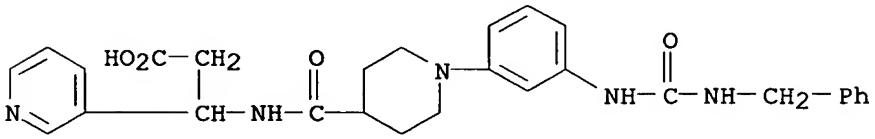


RN 461718-95-6 CAPLUS
CN L-Alanine, N-[(phenylmethoxy)carbonyl]-3-[[[1-[3-[[[(phenylmethyl)amino]carbonyl]amino]phenyl]-4-piperidinyl]carbonyl]amino]- (9CI) (CA INDEX NAME)

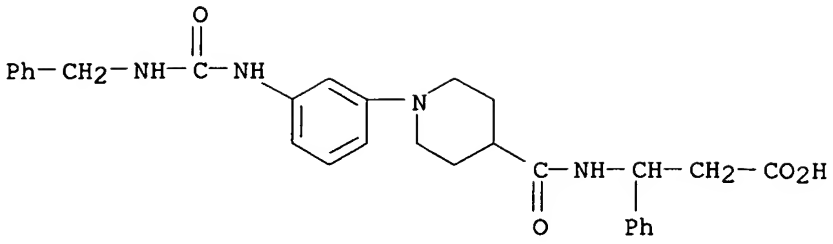
Absolute stereochemistry.



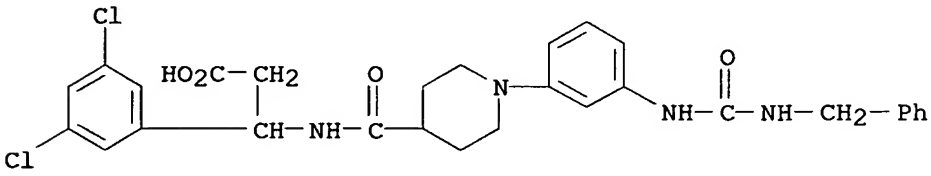
RN 461718-96-7 CAPLUS
CN 3-Pyridinepropanoic acid, β-[[[1-[3-[[[(phenylmethyl)amino]carbonyl]amino]phenyl]-4-piperidinyl]carbonyl]amino]- (9CI) (CA INDEX NAME)



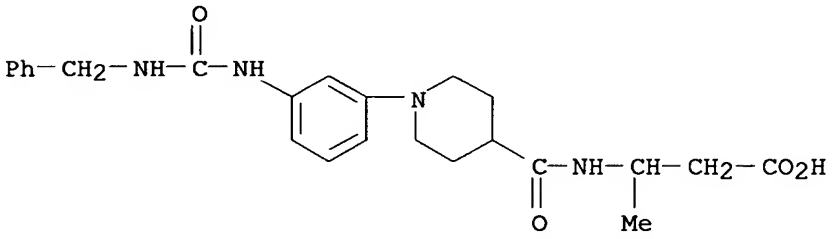
RN 461718-97-8 CAPLUS
CN Benzenepropanoic acid, β-[[[1-[3-[[[(phenylmethyl)amino]carbonyl]amino]phenyl]-4-piperidinyl]carbonyl]amino]- (9CI) (CA INDEX NAME)



RN 461718-98-9 CAPLUS
CN Benzenepropanoic acid, 3,5-dichloro-β-[[[1-[3-[[[(phenylmethyl)amino]carbonyl]amino]phenyl]-4-piperidinyl]carbonyl]amino]- (9CI) (CA INDEX NAME)

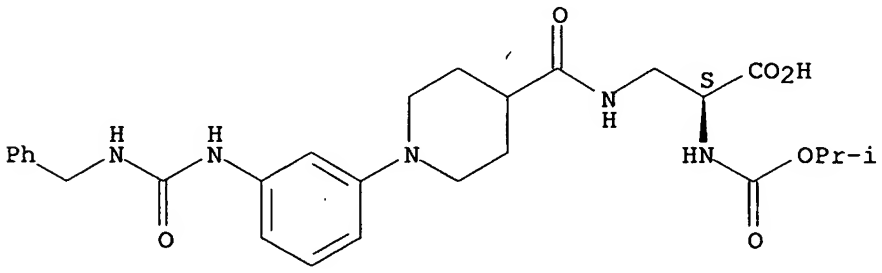


RN 461718-99-0 CAPLUS
CN Butanoic acid, 3-[[[1-[3-[[[(phenylmethyl)amino]carbonyl]amino]phenyl]-4-piperidinyl]carbonyl]amino]- (9CI) (CA INDEX NAME)



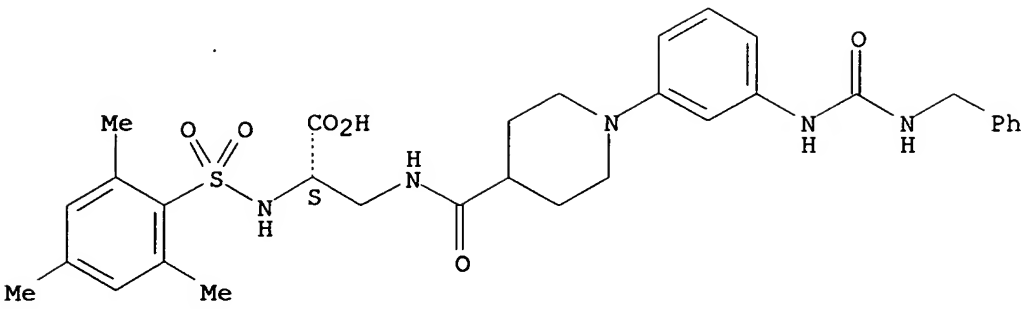
RN 461719-00-6 CAPLUS
CN L-Alanine, N-[(1-methylethoxy)carbonyl]-3-[[[1-[3-[[[(phenylmethyl)amino]carbonyl]amino]phenyl]-4-piperidinyl]carbonyl]amino]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



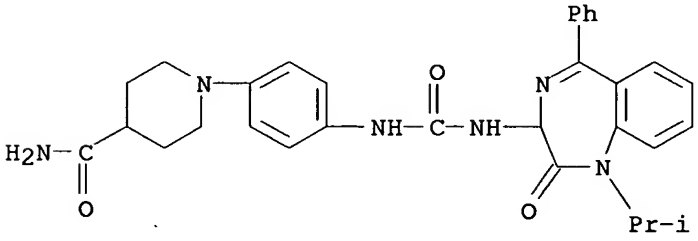
RN 461719-01-7 CAPLUS
CN L-Alanine, 3-[[[1-[3-[[[(phenylmethyl)amino]carbonyl]amino]phenyl]-4-piperidinyl]carbonyl]amino]-N-[(2,4,6-trimethylphenyl)sulfonyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



=> d hitstr 18

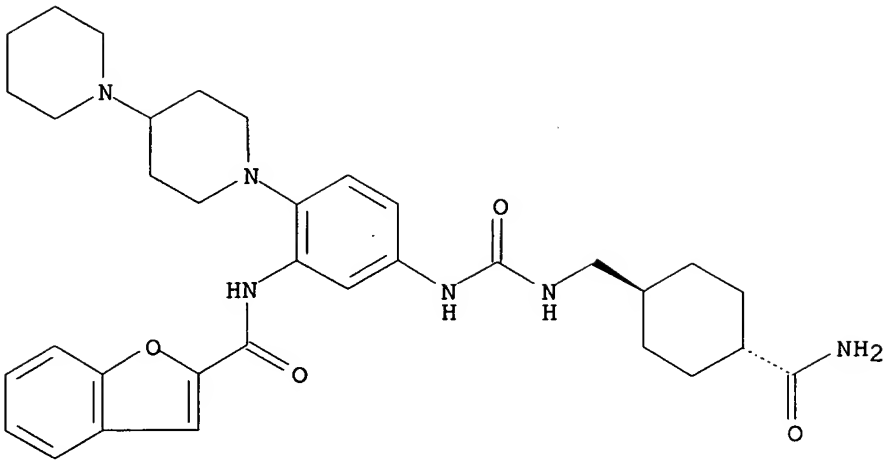
L11 ANSWER 18 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 478055-08-2P
RL: SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(benzodiazepine bradykinin antagonists)
RN 478055-08-2 CAPLUS
CN 4-Piperidinecarboxamide, 1-[4-[[[[2,3-dihydro-1-(1-methylethyl)-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]amino]carbonyl]amino]phenyl]- (9CI) (CA INDEX NAME)



=> d hitstr 17

L11 ANSWER 17 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 488839-58-3P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of substituted amides, sulfonamides and ureas useful for inhibiting kinase activity)
RN 488839-58-3 CAPLUS
CN 2-Benzofurancarboxamide, N-[5-[[[[[trans-4-(aminocarbonyl)cyclohexyl]methyl]amino]carbonyl]amino]-2-[1,4'-bipiperidin]-1'-ylphenyl]- (9CI) (CA INDEX NAME)

Relative stereochemistry.



=> d hitstr 16

L11 ANSWER 16 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 500697-75-6
RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
(method of treating hyperresorptive bone disorders)
RN 500697-75-6 CAPLUS
CN 2-Benzofurancarboxamide, N-[5-[[[[[4-(aminocarbonyl)cyclohexyl]methyl]amino]carbonyl]amino]-2-[1,4'-bipiperidin]-1'-ylphenyl]- (9CI) (CA INDEX NAME)

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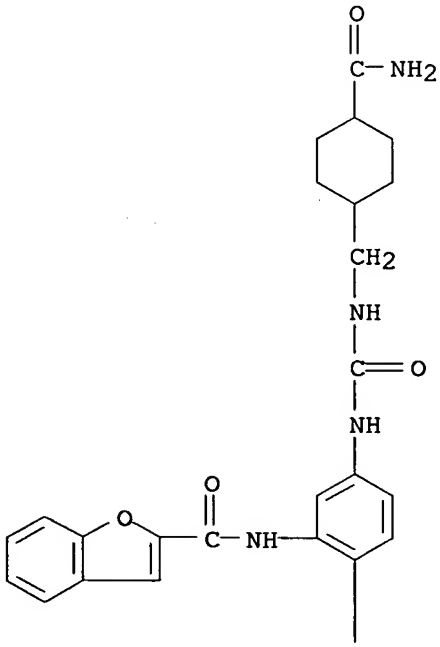
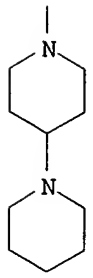
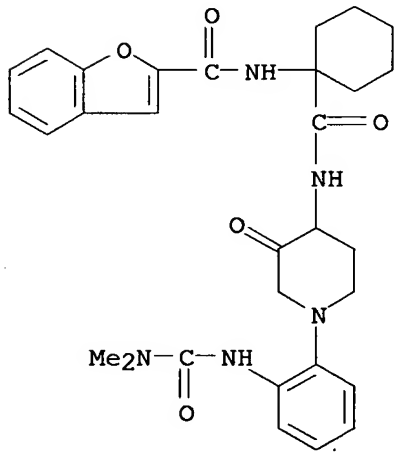


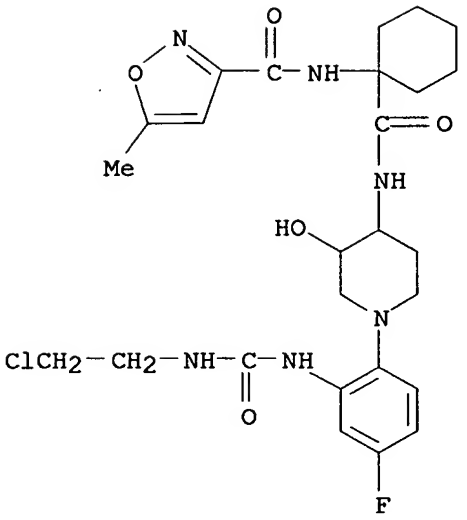


=> d hitstr 15

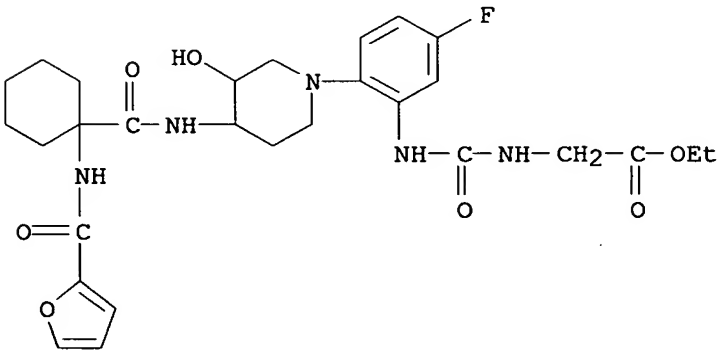
L11 ANSWER 15 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 652171-04-5P, 4-[N-[[1-[N-[(Benzofuran-2-yl)carbonyl]amino]cyclohexyl]carbonyl]amino]-1-[2-(3,3-dimethylureido)phenyl]piperidin-3-one
RL: PAC (Pharmacological activity); PKT (Pharmacokinetics); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation)
(cysteine protease inhibitors; preparation of 4-[[(1-acylaminocyclohexyl)carbonyl]amino]-1-phenylpiperidin-3-ones as cysteine protease inhibitors and processes for their preparation)
RN 652171-04-5 CAPLUS
CN 2-Benzofurancarboxamide, N-[1-[[[1-[2-[[(dimethylamino)carbonyl]amino]phenyl]-3-oxo-4-piperidinyl]amino]carbonyl]cyclohexyl]- (9CI) (CA INDEX NAME)



IT 652171-69-2, 4-[N-[[1-[N-[(5-Methylisoxazol-3-yl)carbonyl]amino]cyclohexyl]carbonyl]amino]-1-[4-fluoro-2-[(2-chloroethylcarbamoyl)amino]phenyl]piperidin-3-ol
RL: RCT (Reactant); RACT (Reactant or reagent)
(preparation of 4-[[(1-acylaminocyclohexyl)carbonyl]amino]-1-phenylpiperidin-3-ones as cysteine protease inhibitors and processes for their preparation)
RN 652171-69-2 CAPLUS
CN 3-Isoxazolecarboxamide, N-[1-[[[1-[2-[[[(2-chloroethyl)amino]carbonyl]amino]-4-fluorophenyl]-3-hydroxy-4-piperidinyl]amino]carbonyl]cyclohexyl]-5-methyl- (9CI) (CA INDEX NAME)

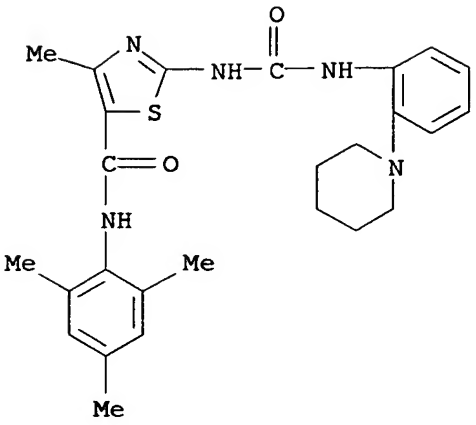


IT 652171-41-0P, 4-[N-[[1-[N-[(Furan-2-yl)carbonyl]amino]cyclohexyl]carbonyl]amino]-1-[4-fluoro-2-[[[(ethoxycarbonyl)methyl]carbamoyl]amino]phenyl]piperidin-3-ol
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation of 4-[[(1-acylaminocyclohexyl)carbonyl]amino]-1-phenylpiperidin-3-ones as cysteine protease inhibitors and processes for their preparation)
RN 652171-41-0 CAPLUS
CN Glycine, N-[[[5-fluoro-2-[4-[[[1-[(2-furanylcarbonyl)amino]cyclohexyl]carbonyl]amino]-3-hydroxy-1-piperidinyl]phenyl]amino]carbonyl]-, ethyl ester (9CI) (CA INDEX NAME)



=> d hitstr 14

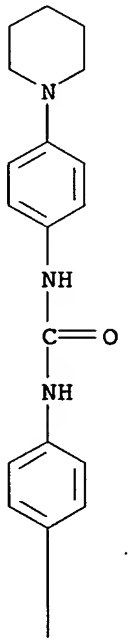
L11 ANSWER 14 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 302960-70-9P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of 5-thiazolecarboxamides as protein tyrosine kinase inhibitors)
RN 302960-70-9 CAPLUS
CN 5-Thiazolecarboxamide, 4-methyl-2-[[[[2-(1-piperidinyl)phenyl]amino]carbonyl]amino]-N-(2,4,6-trimethylphenyl)- (9CI) (CA INDEX NAME)

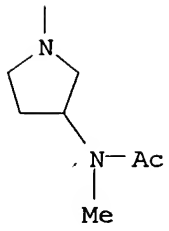


=> d hitstr 13

L11 ANSWER 13 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 748166-67-8P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of N-arylheterocycles as MCH antagonists)
RN 748166-67-8 CAPLUS
CN Acetamide, N-methyl-N-[1-[4-[[[[4-(1-piperidinyl)phenyl]amino]carbonyl]amino]phenyl]-3-pyrrolidinyl]- (9CI) (CA INDEX NAME)

PAGE 1-A

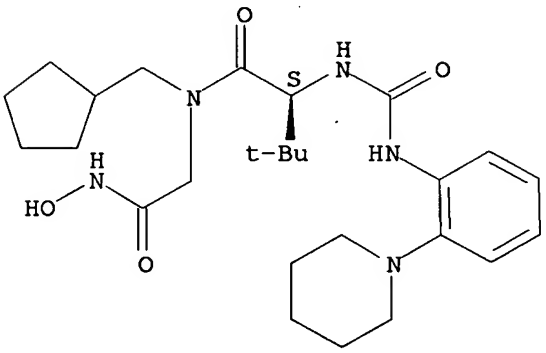




=> d hitstr 12

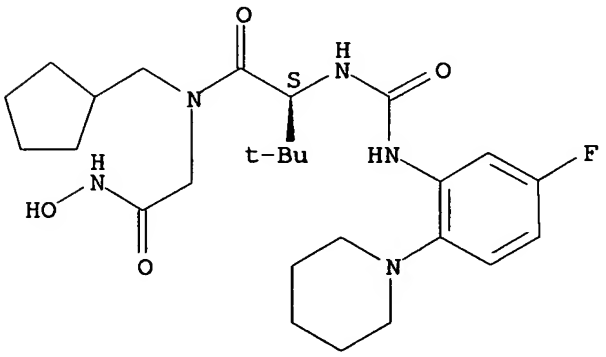
L11 ANSWER 12 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 775320-11-1P 775320-17-7P 775320-20-2P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(reaction of esters such as phenylureidobutyrylaminoacetic acid esters with hydroxylamine)
RN 775320-11-1 CAPLUS
CN Glycinamide, 3-methyl-N-[[[2-(1-piperidinyl)phenyl]amino]carbonyl]-L-valyl-N2-(cyclopentylmethyl)-N-hydroxy- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



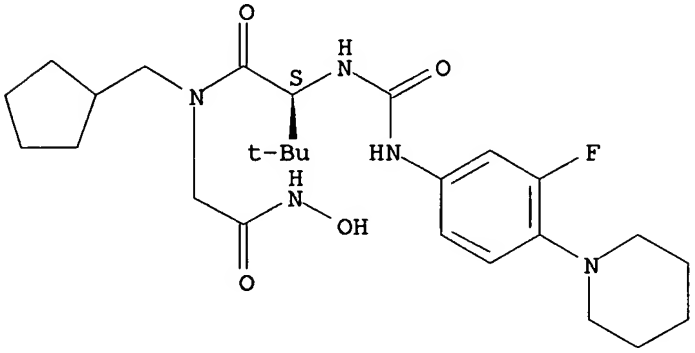
RN 775320-17-7 CAPLUS
CN Glycinamide, N-[[[5-fluoro-2-(1-piperidinyl)phenyl]amino]carbonyl]-3-methyl-L-valyl-N2-(cyclopentylmethyl)-N-hydroxy- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



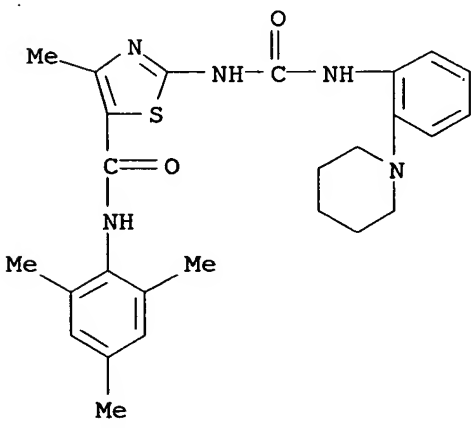
RN 775320-20-2 CAPLUS
CN Glycinamide, N-[[[3-fluoro-4-(1-piperidinyl)phenyl]amino]carbonyl]-3-methyl-L-valyl-N2-(cyclopentylmethyl)-N-hydroxy- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



=> d hitstr 11

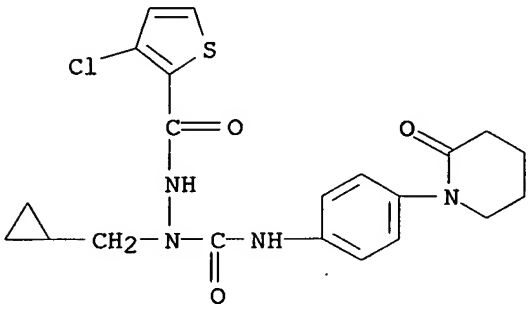
L11 ANSWER 11 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 302960-70-9
RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
(IGF1 receptor inhibitors with addnl. kinase inhibitors for synergistic treatment of cancer)
RN 302960-70-9 CAPLUS
CN 5-Thiazolecarboxamide, 4-methyl-2-[[[[2-(1-piperidinyl)phenyl]amino]carbonyl]amino]-N-(2,4,6-trimethylphenyl)- (9CI) (CA INDEX NAME)



=> d hitstr 9

L11 ANSWER 9 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 808732-16-3P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of aroylsemicarbazides as factor Xa inhibitors for the

treatment of thromboembolic diseases)
RN 808732-16-3 CAPLUS
CN 2-Thiophenecarboxylic acid, 3-chloro-, 2-(cyclopropylmethyl)-2-[[[4-(2-oxo-1-piperidinyl)phenyl]amino]carbonyl]hydrazide (9CI) (CA INDEX NAME)

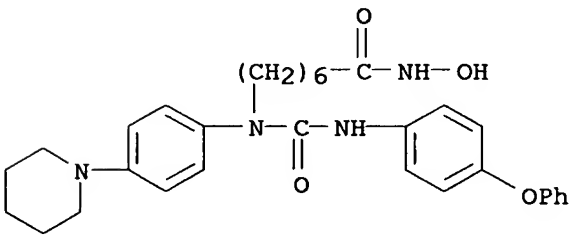


=> d hitstr 8

L11 ANSWER 8 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 811796-03-9P 811796-05-1P 811796-07-3P
811796-09-5P 811796-11-9P 811796-13-1P
811796-15-3P 811796-17-5P 811796-19-7P
811796-52-8P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of N-hydroxy-7-(arylamino)heptanamide derivs. and antitumor activity)
RN 811796-03-9 CAPLUS
CN Heptanamide, N-hydroxy-7-[[[(4-phenoxyphenyl)amino]carbonyl][4-(1-piperidinyl)phenyl]amino]-, mono(trifluoroacetate) (salt) (9CI) (CA INDEX NAME)

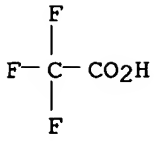
CM 1

CRN 811796-02-8
CMF C31 H38 N4 O4



CM 2

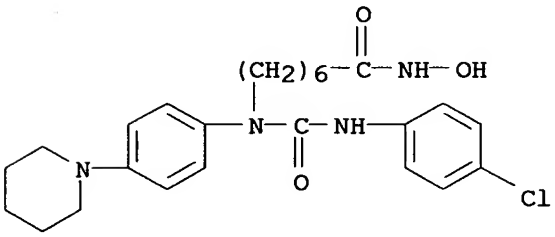
CRN 76-05-1
CMF C2 H F3 O2



RN 811796-05-1 CAPLUS
CN Heptanamide, 7-[[[(4-chlorophenyl)amino]carbonyl][4-(1-piperidinyl)phenyl]amino]-N-hydroxy-, mono(trifluoroacetate) (salt) (9CI) (CA INDEX NAME)

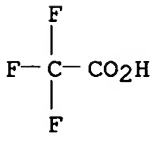
CM 1

CRN 811796-04-0
CMF C25 H33 Cl N4 O3



CM 2

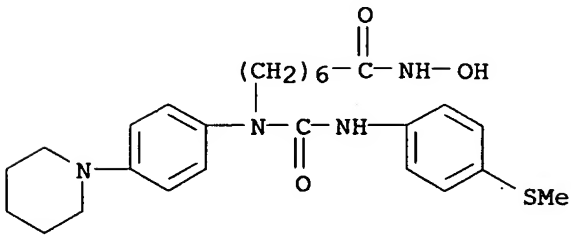
CRN 76-05-1
CMF C2 H F3 O2



RN 811796-07-3 CAPLUS
CN Heptanamide, N-hydroxy-7-[[[[4-(methylthio)phenyl]amino]carbonyl][4-(1-piperidinyl)phenyl]amino]-, mono(trifluoroacetate) (salt) (9CI) (CA INDEX NAME)

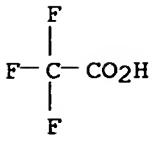
CM 1

CRN 811796-06-2
CMF C26 H36 N4 O3 S



CM 2

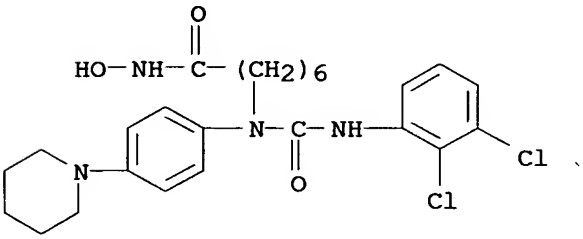
CRN 76-05-1
CMF C2 H F3 O2



RN 811796-09-5 CAPLUS
CN Heptanamide, 7-[[[(2,3-dichlorophenyl)amino]carbonyl][4-(1-piperidinyl)phenyl]amino]-N-hydroxy-, mono(trifluoroacetate) (salt) (9CI) (CA INDEX NAME)

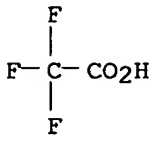
CM 1

CRN 811796-08-4
CMF C25 H32 Cl2 N4 O3



CM 2

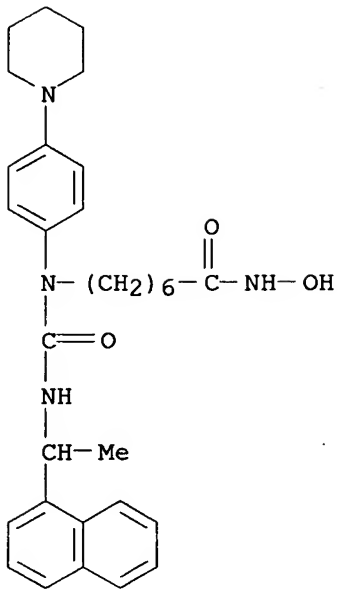
CRN 76-05-1
CMF C2 H F3 O2



RN 811796-11-9 CAPLUS
CN Heptanamide, N-hydroxy-7-[[[[1-(1-naphthalenyl)ethyl]amino]carbonyl][4-(1-piperidinyl)phenyl]amino]-, mono(trifluoroacetate) (salt) (9CI) (CA INDEX NAME)

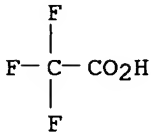
CM 1

CRN 811796-10-8
CMF C31 H40 N4 O3



CM 2

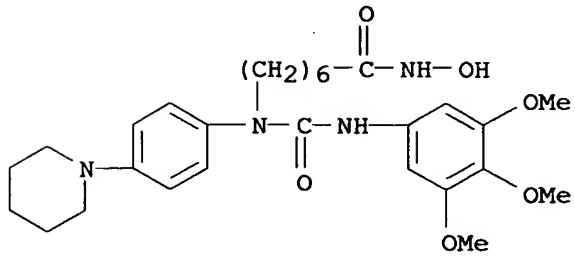
CRN 76-05-1
CMF C2 H F3 O2



RN 811796-13-1 CAPLUS
CN Heptanamide, N-hydroxy-7-[[4-(1-piperidinyl)phenyl][[(3,4,5-trimethoxyphenyl)amino]carbonyl]amino]-, mono(trifluoroacetate) (salt) (9CI) (CA INDEX NAME)

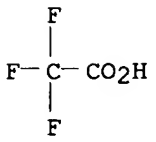
CM 1

CRN 811796-12-0
CMF C28 H40 N4 O6



CM 2

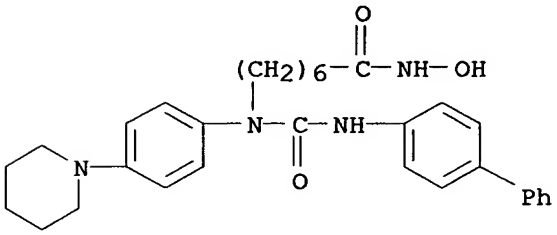
CRN 76-05-1
CMF C2 H F3 O2



RN 811796-15-3 CAPLUS
CN Heptanamide, 7-[[([1,1'-biphenyl]-4-ylamino)carbonyl][4-(1-piperidinyl)phenyl]amino]-N-hydroxy-, mono(trifluoroacetate) (salt) (9CI) (CA INDEX NAME)

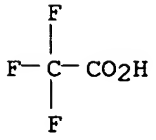
CM 1

CRN 811796-14-2
CMF C31 H38 N4 O3



CM 2

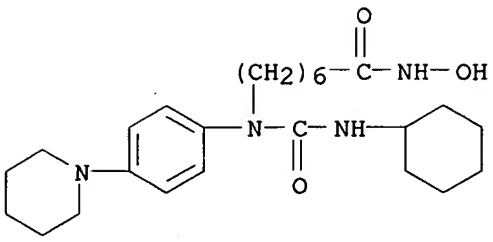
CRN 76-05-1
CMF C2 H F3 O2



RN 811796-17-5 CAPLUS
CN Heptanamide, 7-[[(cyclohexylamino)carbonyl][4-(1-piperidinyl)phenyl]amino]-N-hydroxy-, mono(trifluoroacetate) (salt) (9CI) (CA INDEX NAME)

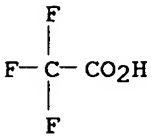
CM 1

CRN 811796-16-4
CMF C25 H40 N4 O3



CM 2

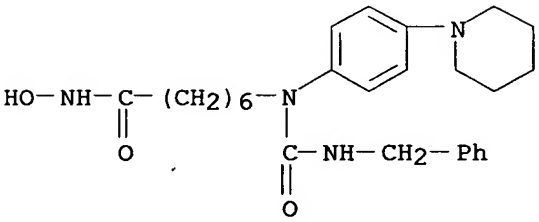
CRN 76-05-1
CMF C2 H F3 O2



RN 811796-19-7 CAPLUS
CN Heptanamide, N-hydroxy-7-[[[(phenylmethyl)amino]carbonyl][4-(1-piperidinyl)phenyl]amino]-, mono(trifluoroacetate) (salt) (9CI) (CA INDEX NAME)

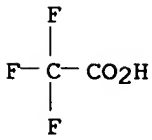
CM 1

CRN 811796-18-6
CMF C26 H36 N4 O3



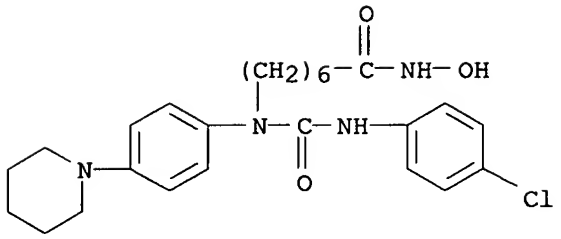
CM 2

CRN 76-05-1
CMF C2 H F3 O2



RN 811796-52-8 CAPLUS
CN Heptanamide, 7-[[[(4-chlorophenyl)amino]carbonyl][4-(1-piperidinyl)phenyl]amino]-N-hydroxy-, monohydrochloride (9CI) (CA INDEX

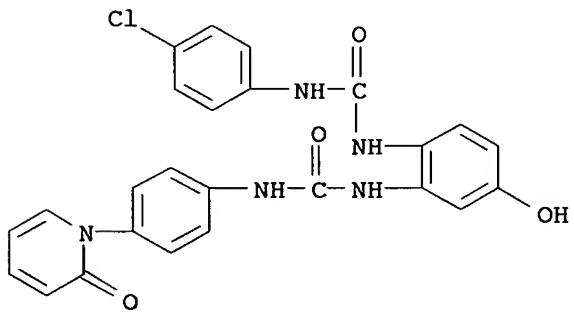
NAME)



● HCl

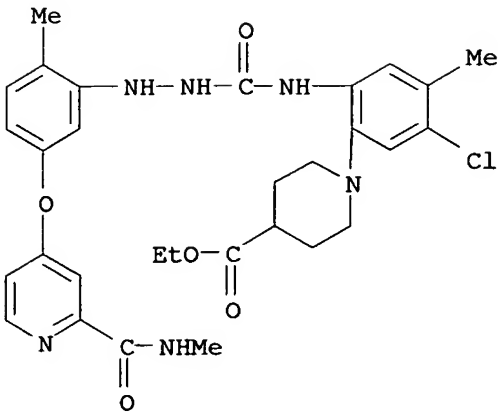
=> d hitstr 7

L11 ANSWER 7 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 862014-74-2P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of heterocyclic urea derivs. as coagulation factor Xa inhibitors)
RN 862014-74-2 CAPLUS
CN Urea, N-[2-[[[(4-chlorophenyl)amino]carbonyl]amino]-5-hydroxyphenyl]-N'-[4-(2-oxo-1(2H)-pyridinyl)phenyl]- (9CI) (CA INDEX NAME)



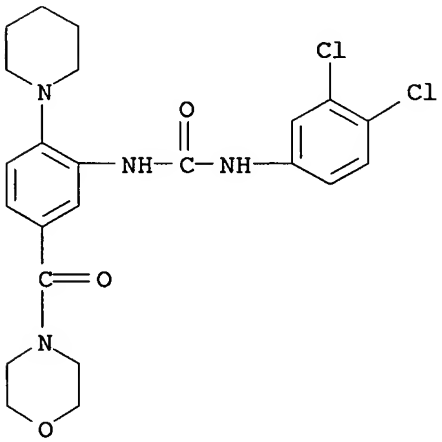
=> d hitstr 6

L11 ANSWER 6 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 864271-94-3P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(drug candidates; preparation of aryl semicarbazides derivs. as inhibitors of raf-kinases, Tie-kinases, PDGFR-kinases and VEGFR-kinases)
RN 864271-94-3 CAPLUS
CN 4-Piperidinecarboxylic acid, 1-[5-chloro-4-methyl-2-[[[2-[2-methyl-5-[[2-[(methylamino)carbonyl]-4-pyridinyl]oxy]phenyl]hydrazino]carbonyl]amino]phenyl]-, ethyl ester (9CI) (CA INDEX NAME)

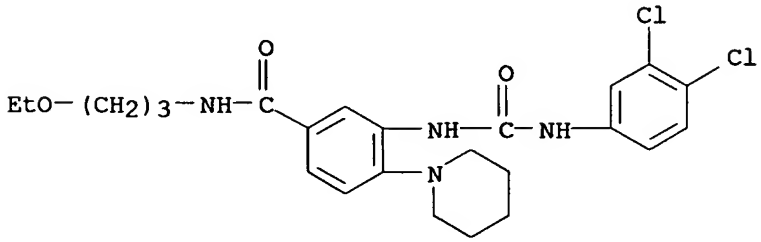


=> d hitstr 5

L11 ANSWER 5 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 898197-27-8P 898197-83-6P
RL: SPN (Synthetic preparation); PREP (Preparation)
(liquid-phase parallel synthesis of 3-substituted 4-aminobenzamides)
RN 898197-27-8 CAPLUS
CN INDEX NAME NOT YET ASSIGNED

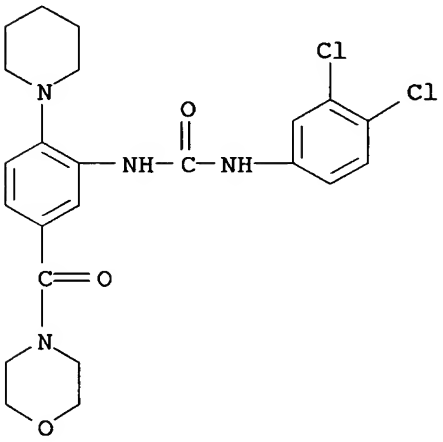


RN 898197-83-6 CAPLUS
CN INDEX NAME NOT YET ASSIGNED

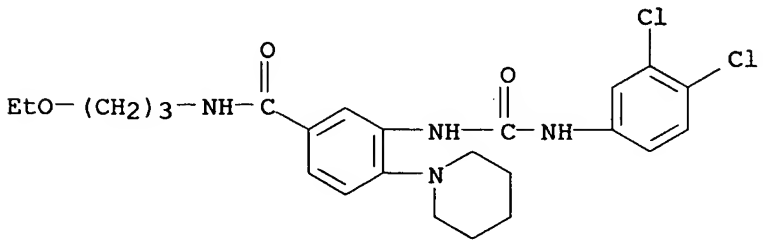


=> d bib hitstr 5

L11 ANSWER 5 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2005:1039511 CAPLUS
DN 145:145355
TI Liquid-phase parallel synthesis of substituted 4-aminobenzamides
AU Trifilenkov, A. S.; Il'in, A. P.; Kravchenko, D. V.; Dorogov, M. V.; Blyumina, M. V.; Ivashchenko, A. V.
CS Yarosl. Gos. Pedagog. Univ. im. K. D. Ushinskogo, Yaroslavl, Russia
SO Izvestiya Vysshikh Uchebnykh Zavedenii, Khimiya i Khimicheskaya Tekhnologiya (2005), 48(5), 137-144
CODEN: IVUKAR; ISSN: 0579-2991
PB Ivanovskii Gosudarstvennyi Khimiko-Tekhnologicheskii Universitet
DT Journal
LA Russian
OS CASREACT 145:145355
IT 898197-27-8P 898197-83-6P
RL: SPN (Synthetic preparation); PREP (Preparation)
(liquid-phase parallel synthesis of 3-substituted 4-aminobenzamides)
RN 898197-27-8 CAPLUS
CN INDEX NAME NOT YET ASSIGNED



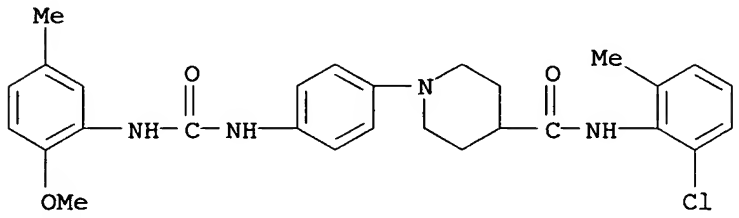
RN 898197-83-6 CAPLUS
CN INDEX NAME NOT YET ASSIGNED



=> d hitstr 4

L11 ANSWER 4 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN

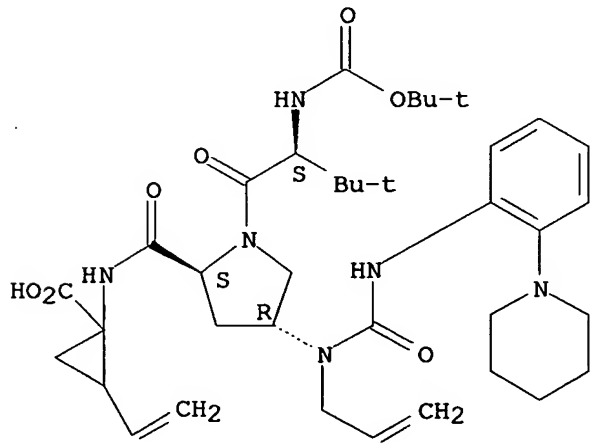
IT 873453-50-0P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of urea derivs. as acyl-CoA:diacylglycerol acyltransferase (DGAT) inhibitors and fat absorption inhibitors)
RN 873453-50-0 CAPLUS
CN 4-Piperidinecarboxamide, N-(2-chloro-6-methylphenyl)-1-[4-[[[(2-methoxy-5-methylphenyl)amino]carbonyl]amino]phenyl]- (9CI) (CA INDEX NAME)



=> d hitstr 3

L11 ANSWER 3 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 877068-71-8P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of tripeptides bearing a cyclopropyl moiety and phosphorous groups as antiviral compds.)
RN 877068-71-8 CAPLUS
CN Cyclopropanecarboxylic acid, N-[(1,1-dimethylethoxy)carbonyl]-3-methyl-L-valyl-(4R)-4-[[[[2-(1-piperidinyl)phenyl]amino]carbonyl]-2-propenylamino]-L-prolyl-1-amino-2-ethenyl- (9CI) (CA INDEX NAME)

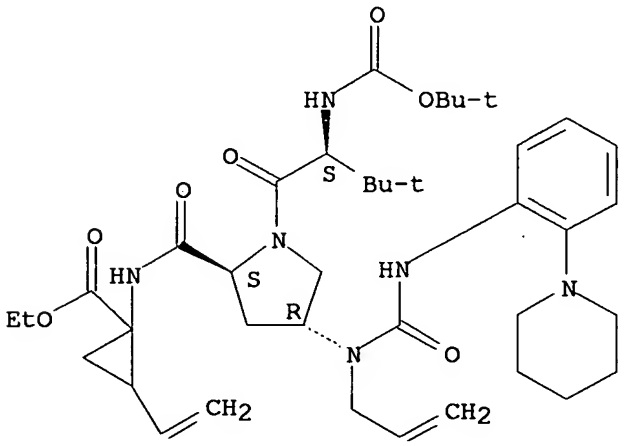
Absolute stereochemistry.



IT 877069-64-2P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation of tripeptides bearing a cyclopropyl moiety and phosphorous groups as antiviral compds.)

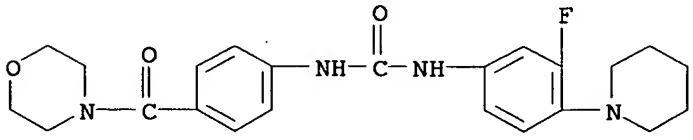
RN 877069-64-2 CAPLUS
CN Cyclopropanecarboxylic acid, N-[(1,1-dimethylethoxy)carbonyl]-3-methyl-L-valyl-(4R)-4-[[[[2-(1-piperidinyl)phenyl]amino]carbonyl]-2-propenylamino]-L-prolyl-1-amino-2-ethenyl-, ethyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.

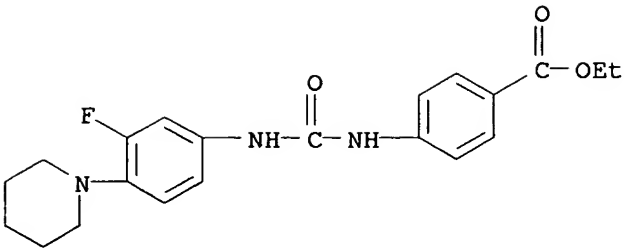


=> d hitstr 2

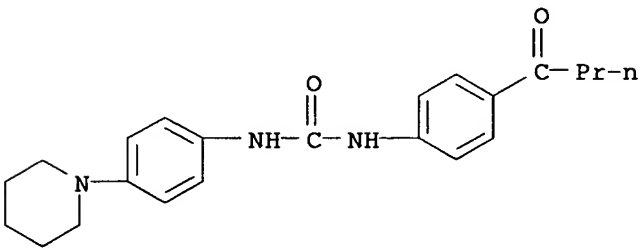
L11 ANSWER 2 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 877202-82-9P, 1-[3-Fluoro-4-(piperidin-1-yl)phenyl]-3-[4-[(morpholin-4-yl)carbonyl]phenyl]urea 877202-84-1P, 4-[3-[3-Fluoro-4-(piperidin-1-yl)phenyl]ureido]benzoic acid ethyl ester 877203-42-4P, 1-(4-Butyrylphenyl)-3-[4-(piperidin-1-yl)phenyl]urea
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(drug candidate; preparation of aryl urea derivs. as CB1 receptor modulators)
RN 877202-82-9 CAPLUS
CN Morpholine, 4-[4-[[[[3-fluoro-4-(1-piperidinyl)phenyl]amino]carbonyl]amino]benzoyl]- (9CI) (CA INDEX NAME)



RN 877202-84-1 CAPLUS
CN Benzoic acid, 4-[[[[3-fluoro-4-(1-piperidinyl)phenyl]amino]carbonyl]amino]-, ethyl ester (9CI) (CA INDEX NAME)



RN 877203-42-4 CAPLUS
CN Urea, N-[4-(1-oxobutyl)phenyl]-N'-[4-(1-piperidinyl)phenyl]- (9CI) (CA INDEX NAME)



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L11 ANSWER 2 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2006:164439 CAPLUS
DN 144:253908
TI Preparation of aryl urea derivatives as CB1 cannabinoid receptor modulators
IN Bloxham, Jason; Fyfe, Matthew Colin Thor; Horswill, James; Jeevaratnam, Revathy Perpetua; Keily, John; Procter, Martin James; Schofield, Karen Lesley; Shaaban, Salam; Swain, Simon Andrew; Wong-Kai-In, Philippe
PA Prosidion Limited, UK
SO PCT Int. Appl., 66 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2006018662	A2	20060223	WO 2005-GB50131	20050816
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
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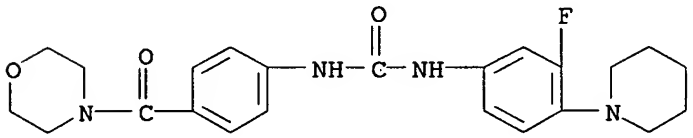
PRAI	US 2004-602268P	P	20040816		

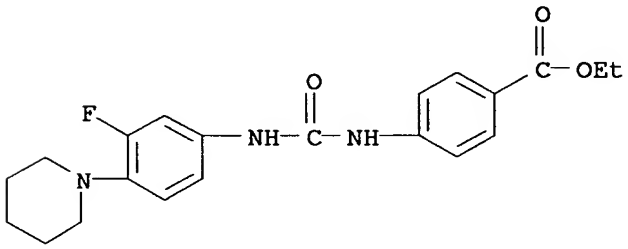
OS MARPAT 144:253908

IT 877202-82-9P, 1-[3-Fluoro-4-(piperidin-1-yl)phenyl]-3-[4-[(morpholin-4-yl)carbonyl]phenyl]urea 877202-84-1P, 4-[3-[3-Fluoro-4-(piperidin-1-yl)phenyl]ureido]benzoic acid ethyl ester 877203-42-4P, 1-(4-Butyrylphenyl)-3-[4-(piperidin-1-yl)phenyl]urea
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(drug candidate; preparation of aryl urea derivs. as CB1 receptor modulators)

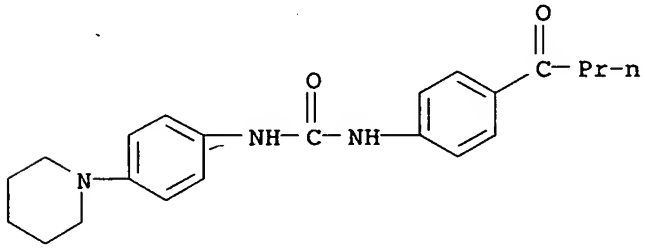
RN 877202-82-9 CAPLUS
CN Morpholine, 4-[4-[[[[3-fluoro-4-(1-piperidinyl)phenyl]amino]carbonyl]amino]benzoyl]- (9CI) (CA INDEX NAME)



RN 877202-84-1 CAPLUS
CN Benzoic acid, 4-[[[[3-fluoro-4-(1-piperidinyl)phenyl]amino]carbonyl]amino]-, ethyl ester (9CI) (CA INDEX NAME)



RN 877203-42-4 CAPLUS
CN Urea, N-[4-(1-oxobutyl)phenyl]-N'-[4-(1-piperidinyl)phenyl]- (9CI) (CA INDEX NAME)



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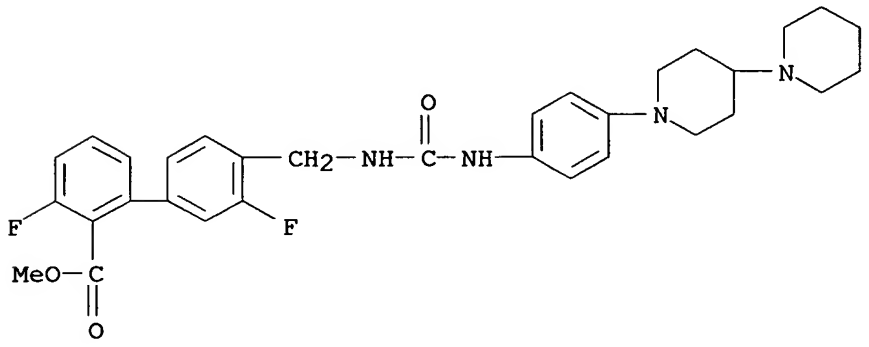
L11 ANSWER 1 OF 37 CAPLUS COPYRIGHT 2006 ACS on STN
IT 887142-70-3 887142-71-4 887142-73-6
887142-75-8 887142-78-1 887142-79-2
887142-80-5 887142-82-7 887142-83-8
RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(substituted biaryl-carboxylate derivs. as bradykinin B1 antagonists or inverse agonists useful in the treatment of pain and inflammation)

RN 887142-70-3 CAPLUS

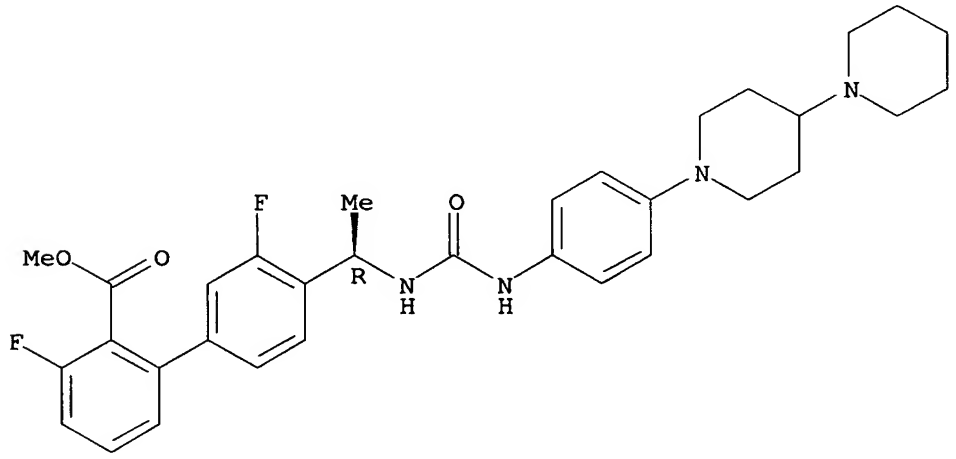
CN [1,1'-Biphenyl]-2-carboxylic acid, 4'-[[[[(4-[1,4'-bipiperidin]-1'-ylphenyl)amino]carbonyl]amino]methyl]-3,3'-difluoro-, methyl ester (9CI) (CA INDEX NAME)



RN 887142-71-4 CAPLUS

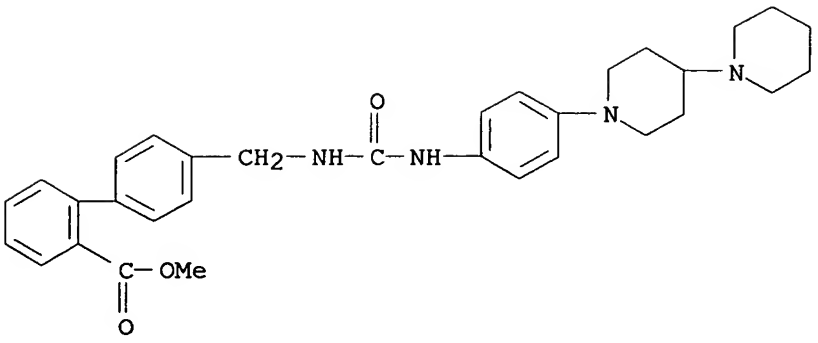
CN [1,1'-Biphenyl]-2-carboxylic acid, 4'-[(1R)-1-[[[(4-[1,4'-bipiperidin]-1'-ylphenyl)amino]carbonyl]amino]ethyl]-3,3'-difluoro-, methyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.

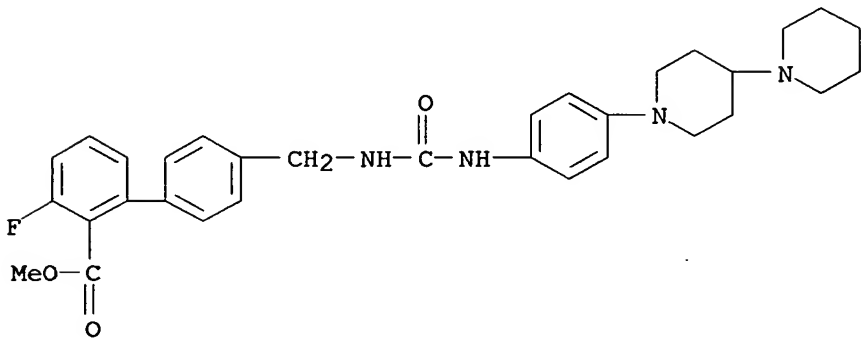


RN 887142-73-6 CAPLUS

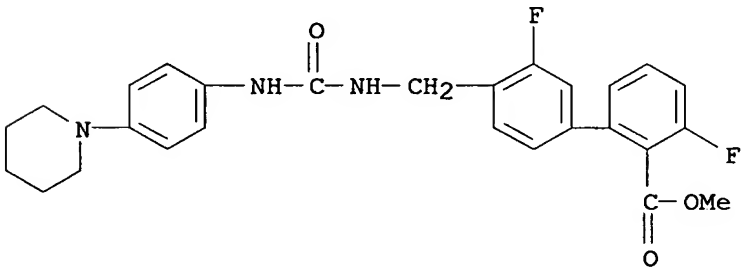
CN [1,1'-Biphenyl]-2-carboxylic acid, 4'-[[[[(4-[1,4'-bipiperidin]-1'-ylphenyl)amino]carbonyl]amino]methyl]-, methyl ester (9CI) (CA INDEX NAME)



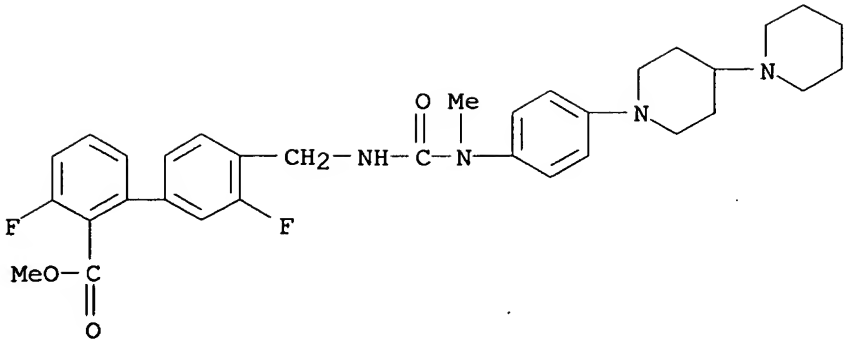
RN 887142-75-8 CAPLUS
CN [1,1'-Biphenyl]-2-carboxylic acid, 4'-[[[[(4-[1,4'-bipiperidin]-1'-ylphenyl)amino]carbonyl]amino]methyl]-3-fluoro-, methyl ester (9CI) (CA INDEX NAME)



RN 887142-78-1 CAPLUS
CN [1,1'-Biphenyl]-2-carboxylic acid, 3,3'-difluoro-4'-[[[[[4-(1-piperidinyl)phenyl]amino]carbonyl]amino]methyl]-, methyl ester (9CI) (CA INDEX NAME)

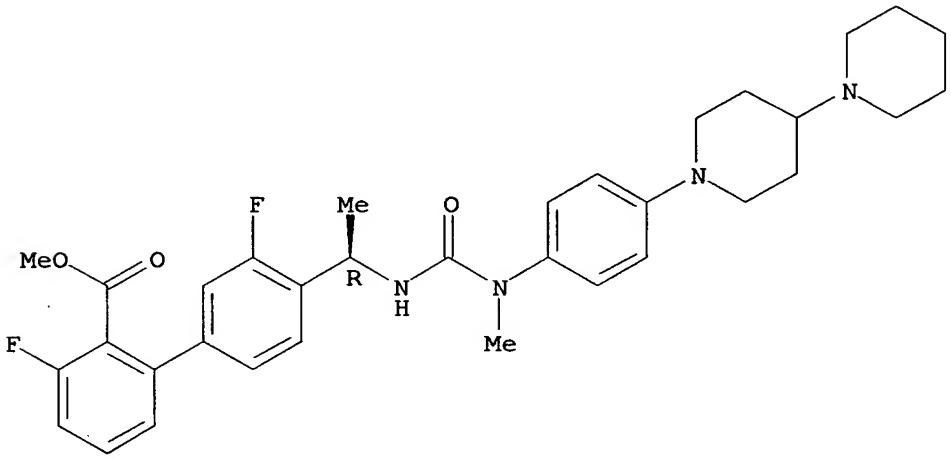


RN 887142-79-2 CAPLUS
CN [1,1'-Biphenyl]-2-carboxylic acid, 4'-[[[[(4-[1,4'-bipiperidin]-1'-ylphenyl)methylamino]carbonyl]amino]methyl]-3,3'-difluoro-, methyl ester (9CI) (CA INDEX NAME)

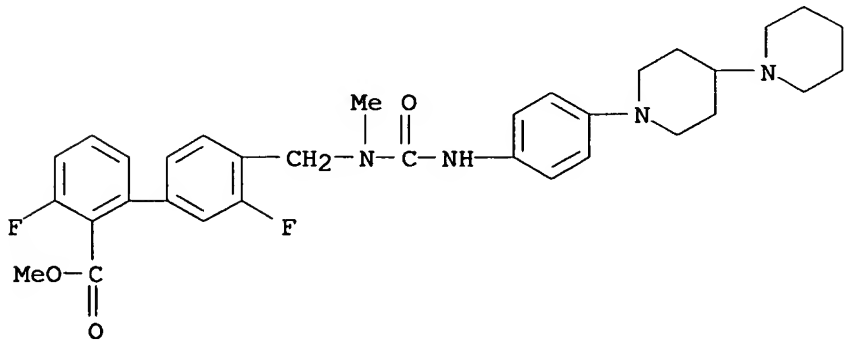


RN 887142-80-5 CAPLUS
CN [1,1'-Biphenyl]-2-carboxylic acid, 4'-[(1R)-1-[[[(4-[1,4'-bipiperidin]-1'-ylphenyl)methylamino]carbonyl]amino]ethyl]-3,3'-difluoro-, methyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.



RN 887142-82-7 CAPLUS
CN [1,1'-Biphenyl]-2-carboxylic acid, 4'-[[[[(4-[1,4'-bipiperidin]-1'-ylphenyl)amino]carbonyl]methylamino]methyl]-3,3'-difluoro-, methyl ester (9CI) (CA INDEX NAME)



RN 887142-83-8 CAPLUS
CN [1,1'-Biphenyl]-2-carboxylic acid, 3,3'-difluoro-4'-[[[[[4-[4-(4-pyridinyl)-1-piperidinyl]phenyl]amino]carbonyl]amino]methyl]-, methyl

ester (9CI) (CA INDEX NAME)